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- (71) Applicant (*for all designated States except US*): HUMAN GENOME SCIENCES, INC. [US/US]; 9410 Key West Avenue, Rockville, MD 20850 (US).
- (72) Inventors; and
- (75) Inventors/Applicants (*for US only*): ROSEN, Craig, A. [US/US]; 22400 Rolling Hill Lane, Laytonsville, MD 20882 (US). BIRSE, Charles, E. [GB/US]; 13822 Saddleview Drive, North Potomac, MD 20878 (US).
- (74) Agents: WALES, Michele, M. et al.; Human Genome Sciences, Inc., 9410 Key West Avenue, Rockville, MD 20850 (US).
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(54) Title: NUCLEIC ACIDS, PROTEINS, AND ANTIBODIES

(57) Abstract: The present invention relates to novel proteins. More specifically, isolated nucleic acid molecules are provided encoding novel polypeptides. Novel polypeptides and antibodies that bind to these polypeptides are provided. Also provided are vectors, host cells, and recombinant and synthetic methods for producing human polynucleotides and/or polypeptides, and antibodies. The invention further relates to diagnostic and therapeutic methods useful for diagnosing, treating, preventing and/or prognosing disorders related to these novel polypeptides. The invention further relates to screening methods for identifying agonists and antagonists of polynucleotides and polypeptides of the invention. The present invention further relates to methods and/or compositions for inhibiting or enhancing the production and function of the polypeptides of the present invention.

Nucleic Acids, Proteins, and Antibodies

Field of the Invention

[0001] The present invention relates to novel proteins. More specifically, isolated nucleic acid molecules are provided encoding novel polypeptides. Novel polypeptides and antibodies that bind to these polypeptides are provided. Also provided are vectors, host cells, and recombinant and synthetic methods for producing human polynucleotides and/or polypeptides, and antibodies. The invention further relates to diagnostic and therapeutic methods useful for diagnosing, treating, preventing and/or prognosing disorders related to these novel polypeptides. The invention further relates to screening methods for identifying agonists and antagonists of polynucleotides and polypeptides of the invention. The present invention further relates to methods and/or compositions for inhibiting or enhancing the production and function of the polypeptides of the present invention.

Background of the Invention

[0002] The invention embodies novel CD-like molecules. CD molecules are membrane proteins that were first identified as phenotypic markers for different lymphocyte

populations. CD stands for "cluster of differentiation" and refers to a molecule recognized by a "cluster" of monoclonal antibodies that can be used to identify the lineage or distinguish one class of lymphocytes from another. CD proteins were first used for subclassifying T cells, but different CD molecules recognized by specific antibodies now serve as useful markers of B cells and other leukocytes that participate in immune and inflammatory responses. A number of CD molecules grouped by their molecular families are discussed below. For further information, see, e.g., *Leukocyte Typing IV. White Cell Differentiation Antigens*, Knapp, Dorken, Gilks et al. (eds.). Oxford, Oxford University Press, 1989; hereby incorporated in its entirety by reference herein.

[0003] **Immunoglobulin superfamily**: The Immunoglobulin (Ig) Superfamily of proteins consists of diverse families of proteins involved in cell-cell interactions, cell-surface recognition, intercellular communication, and immune and inflammatory responses. Members of the gene superfamily are structural homologs, but they do not necessarily share similar functions or proximity in the genome. Most identified members of the Ig superfamily are integral plasma membrane proteins with Ig domains in the extracellular portions, transmembrane domains composed of hydrophobic amino acids, and widely divergent cytoplasmic tails with little or no homology to one another or to previously identified signal-transduction structures. Members of this superfamily of proteins include molecules such as Cell Adhesion Molecules (CAMs), CD1, CD2, CD3d, CD4, CD7, CD8a, CD16a, CD19, CD22, CD23, CD28, CD31, CD33, CD47, CD48, CD50, CD54, CD56, CD58, CD64, CD66a, CD79a, CD80, CD83, CD84, CD86, CD89 (Fc receptor), CD90, CD96, CD100, CD101, CD102, CD106, CD115, CD117, CD121a, CD146, CD147, CD152, CD158a, CD166, and T-cell receptor, for example.

[0004] Neural CAMs are thought to play important roles in axonal growth and guidance. Intercellular adhesion molecules (ICAMs), Vascular adhesion molecule-1 (VCAM-1), and Mucosal addressin cell adhesion molecule-1 (MAdCAM-1) are all thought to be involved in binding to integrins of leukocyte surfaces, thus facilitating the immune response to infection and injury (Wang, J., and T.A. Springer, *Immun. Reviews*, 163:197-215 (1998)).

[0005] CD1 molecules have been demonstrated to restrict T cell responses to non-peptide lipid and glycolipid antigens. CD2 functions as an intercellular adhesion molecule and is involved with signal transduction. CD4 and CD8 are T cell surface glycoproteins

that serve as accessory molecules by facilitating interactions of T cells with APCs or CTL target cells. CD19 is a critical signal transduction molecule that regulates B lymphocyte development, activation, and differentiation. CD23 is involved in the regulation of IgE synthesis. CD54 reacts with CD11a/CD18 or CD11b/CD18 resulting in immune reaction and/or inflammation. CD58 mediates the adhesion between killer and target cells, antigen presenting cells and T cells, and thymocytes and thymic epithelial cells. CD64 is involved in antigen capture for presentation to T cells and receptor-mediated endocytosis of IgG-antigen complexes. CD86 and CD80 acts as ligands for the T cell stimulation. CD115 (CSF-1R) is the receptor for CSF-1 and mediates all of the biological effects of this cytokine.

[0006] These molecules function in cell-cell interactions, cell-surface recognition, intercellular communication, and immune and inflammatory responses, related molecules would be predicted to have similar biological activities.

[0007] Further members of the immunoglobulin superfamily, which also fall into the cytokine receptor family, include, for example, CD114, CD116, CD122, CD123, CD124, CDw125, CD126, CD127, CD130, CDw131, CD132, CDw199.

[0008] These molecules function as cytokine receptors, related molecules would be predicted to have a similar biological activity. Such as, for example, CD132 is the receptor for IL-2, IL-4, IL-7, IL-9 and IL-15. Mutations in humans causes XSCID, a disease characterized by the absence of T and NK cells. CDw131 is a key signal transducing molecule of the IL-3, GM-CSF and IL-5 receptors. CD127 is the receptor for interleukin-7. CD124 is a receptor subunit for Interleukin-4 and 13.

[0009] **Integrin assemblies:** The integrin gene family consists of homologous genes that encode molecules which promote cell-cell or cell-matrix interactions. The integrin family of heterodimeric receptors provides structural links between the extracellular environment and the cytoskeleton-signaling network and thus helps to regulate cell migration, differentiation, cell cycle progression, apoptosis, phagocytosis, ECM assembly, and metalloproteinase activity. All integrins are heterodimeric cell surface proteins composed of two noncovalently linked polypeptide chains, alpha and beta. The two chains consist of extracellular and transmembrane segments and cytoplasmic tails 30 to 46 amino acids long. The extracellular domains of the two chains bind to various ligands, including

extracellular matrix glycoproteins, complement components and proteins on the surface of other cells. Many of the integrins bind to Arg-Gly-Asp (R-G-D) sequences in the ligands, but this is not always the case. The cytoplasmic domains of the integrins interact with cytoskeletal components (including vinculin, talin, actin, alpha-actinin, and tropomyosin), and it is hypothesized that the integrins coordinate (i.e., "integrate") the binding of cells to extracellular proteins with cytoskeleton-dependent motility, shape change and phagocytic responses. Members of the integrin superfamily include molecules involved in beta chain assembly such as, for example, CD104, CD61, CD29, CD18; and molecules involved in alpha chain assembly, such as, CD103, CD51, CD49a, CD41, CD11c, CD11b, CD11a, for example.

[0010] Three integrin subfamilies were originally defined on the basis of which of three beta subunits were used to form the heterodimers. The beta1-containing integrins are also called "very late activating" (VLA) molecules. The VLA proteins mediate attachment of leukocytes to extracellular matrices and vascular endothelium.

[0011] The beta2-integrins, also known as the LFA-1 family, play an important role in the adhesion of lymphocytes with other cells, such as accessory cells and vascular endothelium. An autosomal recessive inherited deficiency in LFA-1, Mac-1 and p150,95 proteins, called leukocyte adhesion deficiency (the result of a mutation in the CD18 gene), has been identified in a few families and is characterized by recurrent bacterial and fungal infections, lack of polymorphonuclear leukocyte accumulations and defects in adherence-dependent immune functions.

[0012] These molecules promote cell-cell or cell-matrix interactions, related molecules would be predicted to have similar biological activities.

[0013] **TM4SF:** Cell surface proteins of the tetraspanin superfamily (or TM4SF) are a newly characterized family of proteins which are presumed to span the plasma membrane four times. The function of this family of molecules is poorly understood, but there is evidence that they may be involved in transmembrane signal transduction and regulation of cellular proliferation, development, motility, and tumor cell growth and metastasis, in a number of different cell types. The TM4SF proteins may function as "adaptors" or "molecular facilitators," organizing various cell-surface proteins into large multimeric complexes on the cell surface. The TM4SF proteins contain putative small (20-27 amino

acids) and large (75-130 amino acids) extracellular domains and four putative hydrophobic transmembrane domains. Many TM4SF proteins are thought to be involved in signal transduction. In fact, several TM4SF members have been shown to associate with tyrosine phosphatases. Members of this family include, CD151, CD82, CD81, CD63, CD53, CD37, CD9, for example. Several of these members, including CD151, CD81, CD63 and CD9, are known to interact with integrin $\alpha 3/\beta 1$ integrin complexes.

[0014] A brief overview indicates that some tetraspanins (e.g., CD81, CD82, CD9, CD63) are found in virtually all tissues, whereas others are highly restricted, such as CD37 (B cells) or CD53 (lymphoid and myeloid cells). Many of these proteins have a flair for promiscuous associations with other molecules, including lineage-specific proteins, integrins, and other tetraspanins. In terms of function, they are involved in diverse processes such as cell activation and proliferation, adhesion and motility, differentiation, and cancer. It is proposed that these functions may all relate to their ability to act as "molecular facilitators," grouping specific cell-surface proteins and thus increasing the formation and stability of functional signaling complexes.

[0015] These molecules act as "molecular facilitators," grouping specific cell-surface proteins and thus increasing the formation and stability of functional signaling complexes, related molecules would be predicted to have similar biological activities.

[0016] **Extracellular enzymes:** Several CD molecules have enzymatic activity. Such as, for example, protein kinase activity. Members include, for example, CDw136 (involved in the induction of migration, morphological change, and proliferation in different target cells), and CD42a-d (CD42a-d complex serves as the receptor for von Willebrand factor and thrombin -- the complex mediates adhesion of platelets to subendothelial matrices).

[0017] Other members function as phosphatases, such as, for example, CD148, and CD45 (a tyrosine phosphatase which is critical for T and B cell antigen receptor-mediated activation). Further members function as, ectoenzymes, such as for example, CD157 (ADP-ribosyl cyclase and cyclic ADP-ribose hydrolase activity), CD38 (involved in the metabolism of two calcium messengers cADPR and NAADP), CD26 (cleaves off N-terminal X-Pro or X-Ala dipeptides from polypeptides), and CD39 (ecto-apyrase). Moreover, members function as peptidases, such as, for example, metallopeptidases:

CD13 (catalyzes the removal of single amino acids from the amino terminus of small peptides and probably plays a role in their final digestion), CD156, CD143 (peptidyl dipeptide hydrolase involved in the metabolism of two major vaso-active peptides, angiotensin II and bradykinin), and CD10. Other members function as proteoglycans, for example, CD138, or protease cofactors, for example, CD142.

[0018] These molecules have enzymatic activity, related molecules would be predicted to have similar biological activities.

[0019] **Scavenger Receptor Superfamily:** Scavenger receptors (SRs) have been studied primarily due to their ability to bind and internalize modified lipoproteins, suggesting an important role in foam cell formation and the pathogenesis of atherosclerosis. However, the ability of some SRs to function as pattern recognition receptors through their binding of a wide variety of pathogens indicates a potential role in host defence. The scavenger receptor family of proteins includes: class A (type I and II macrophage scavenger receptors, MARCO), class B (CD36, scavenger receptor class BI), mucinlike (CD68/macrosialin, dSR-CI) and endothelial (LOX-1) receptors. Two motifs have been identified as ligand-binding domains: a charged collagen structure of type I and II receptors, and an immunodominant domain of CD36. These structures can recognize a wide range of negatively charged macromolecules, including oxidized low-density lipoproteins, damaged or apoptotic cells, and pathogenic microorganisms. After binding, these ligands can be either internalized by endocytosis or phagocytosis, or remain at the cell surface and mediate adhesion or lipid transfer through caveolae. Under physiological conditions, scavenger receptors serve to scavenge or clean up cellular debris and other related materials, and they play a role in host defence. In pathological states, they mediate the recruitment, activation and transformation of macrophages and other cells which may be related to the development of atherosclerosis and to disorders caused by the accumulation of denatured materials, such as Alzheimer's disease. Scavenger-receptor class A has been held responsible for the clearance of modified LDL from the blood circulation. Scavenger-receptor BI can facilitate selective uptake of cholesterol esters from HDL. Members of the scavenger receptor superfamily include molecules such as, CD163, CD36, CD68, CD6, and CD5, for example.

[0020] These molecules can recognize a wide range of negatively charged macromolecules, including oxidized low-density lipoproteins, damaged or apoptotic cells, and pathogenic microorganisms, related molecules would be predicted to have similar biological activities.

[0021] **Regulators of Complement Activation Gene:** The C3 convertases of the human complement system are controlled by fluid-phase and membrane proteins in the RCA (regulators of complement activation) family. The RCA members are related structurally (possess approximately 60 amino acid repeating motifs termed short consensus repeats (SCR)), functionally (bind C3b/C4b), and genetically (genes are tightly clustered on chromosome 1 at q3.2). Uncontrolled activation of complement can lead to the formation of the membrane attack complex on self tissues and excessive generation of inflammatory mediators.

[0022] Members of this family include molecules such as, CD46 (cofactor for Factor 1 proteolytic cleavage of C3b and C4b); CD35 (mediates adherence of C4b/C3b coated particles in preparation for phagocytosis); and CD21 (part of a large signal transduction complex involving CD19, CD81 and Leu13), for example. Other molecules, such as, CD59, for example, are also involved in the regulation of complement activation, but contain no SCRs.

[0023] These molecules regulators of complement activation, related molecules would be predicted to have similar biological activities.

[0024] **C-type lectins:** C-type lectins (C-TLs) are a family of carbohydrate-binding proteins intimately involved in diverse processes including vertebrate immune cell signalling and trafficking, activation of innate immunity in both vertebrates and invertebrates, and venom-induced haemostasis. Members of this family include, for example, CD62E, CD62L, CD62P, CD69, CD72, CD94, CD141, and CD161.

[0025] CD69 is a member of the natural killer (NK) cell gene complex family of signal transducing receptors and may be involved in the pathogenesis of such diseases as rheumatoid arthritis, chronic inflammatory liver diseases, mild asthma, and acquired immunodeficiency syndrome. CD62E, L, and P mediate leukocyte rolling on activated endothelium at inflammatory sites. CD94 forms a complex with NKG2-A which is

involved in inhibition of NK cell function. CD141 is critical for the activation of protein C and initiation of the protein C anticoagulant pathway.

[0026] These molecules are involved in diverse processes including vertebrate immune cell signalling and trafficking, activation of innate immunity in both vertebrates and invertebrates, and venom-induced haemostasis, related molecules would be predicted to have similar biological activities.

[0027] **TNFs and TNF receptor families:** Tumor necrosis factors (TNF) alpha and beta are cytokines, which act through TNF receptors to regulate numerous biological processes, including protection against infection and induction of shock and inflammatory disease. The TNF molecules belong to the "TNF-ligand" superfamily, and act together with their receptors or counter-ligands, the "TNF-receptor" superfamily. TNF family members include, for example, CD70, CD153, and CD154. TNF-R family members include, for example, CD27 (mediates the co-stimulatory signal for T and B cell activation), CD30 (involved in TCR mediated cell death), CD40 (central to T dependent responses), CD95 (involved in the mediation of apoptosis inducing signals), CD120a, CD 134, and CDw137 (costimulator of T cell proliferation).

[0028] These molecules are involved in providing protection against infection and induction of shock and inflammatory disease, related molecules would be predicted to have similar biological activities.

[0029] **Mucins:** Mucins are adhesion molecules that play a part in leukocyte migration, homing and cell-cell interactions. Mucins are thought to bind to L-selectin and initiate leukocyte:endothelial cell interactions. Members include, for example, CD34, CD42b (part of the CD42a-d complex which serves as a receptor for the von Willebrand factor and thrombin), CD43, CD99, CD162 (mediates rolling of leukocytes on activated endothelium, on activated platelets and on other leukocytes at inflammatory sites), and CD164.

[0030] These molecules play a part in leukocyte migration, homing and cell-cell interactions, related molecules would be predicted to have similar biological activities.

[0031] **Other types of CD molecules:** Several CD molecules are known to be anchored at the cell surface by glycosyl phosphatidyl inositol. Members include, for example, CD55, CD59, CD73, CD87, and CDw108. Genetic defects in GPI-anchor attachment that cause a reduction or loss of both CD55 and CD59 on erythrocytes produce the symptoms of the disease paroxysmal nocturnal hemoglobinuria (PNH). CD73, like other GPI anchored surface proteins, can mediate costimulatory signals in T cell activation. Very small (e.g., 12-35 amino acids) CD molecules which are formed by displacement of C terminal amino acids by GPI linkage include, for example, CD24 and CD52.

[0032] Other CD molecules are members of receptor families, such as, for example, G-protein coupled receptors (e.g., CD97, and CD88); tyrosine kinase receptors (e.g., CD135, and CD140a); transferrin receptors (e.g., CD71); LDL receptor (e.g., CD91); TGF-beta type III receptors (e.g., CD105); Fc receptors for IgG (e.g., CD32); chemokine receptors (e.g., CDw128a); and viral receptors (e.g., CD155).

[0033] Many cell surface carbohydrates have been given CD designation. These include, for example, CD15s, CD 15, CDw17, CD57, CDw60 (CDw60 antibodies provide costimulatory/comitogenic signals for T cells), CD65, CD65s, CDw75 (cell adhesion, ligand for CD22), CDw76, and CD77 (cross-linking of CD77 induces apoptosis of Burkitt's lymphoma cells).

[0034] Many CD molecules have been assigned to small molecular families. These include, for example, members of the lamp family (e.g., CD107a); members of the sushi family (e.g., CD25, the IL-2 receptor); membrane glycoproteins (e.g., CD165, which is involved in the adhesion between thymocytes and thymic epithelial cells).

[0035] Other CD molecules have not yet been assigned to a molecular family, for example, CDw12; CD14 (functions as a receptor for endotoxin [LPS]); CD20; CD44 and CD44R (may be involved in leukocyte attachment to and rolling on endothelial cells, homing to peripheral lymphoid organs and to sites of inflammation); CD74 (involved in the intracellular sorting of MHC class II molecules); CD85 (may play a role in T cell activation in peripheral blood); CDw92; CDw93; CD98 (involved in the regulation of cellular activation); CDw150 (binding of CDw150 to its ligand enhances proliferation and Ig production by activated B cells; CD139; and CD109.

[0036] These molecules may function as described above, related molecules would be predicted to have similar biological activities.

[0037] CD-like proteins are cell surface molecules which play various functions in the immune system, including cell-cell interaction, regulation of inflammation.

[0038] Further, CD-like proteins with a predominant tissue expression pattern are important targets for targeted drug delivery, tumor-targeted therapy (e.g., including, but not limited to, radioimmunotherapy) antibody mediated attack of diseased tissues or cancers, and immune mediated cytotoxicity.

[0039] The discovery of new CD-like proteins and the polynucleotides encoding these molecules thus satisfies a need in the art by not only providing new compositions useful in the diagnosis, treatment, and prevention of diseases and disorders of the immune system and/or associated with tissues where expression is predominant; but also by providing new targets for immune based therapies.

Summary of the Invention

[0040] The present invention relates to novel proteins. More specifically, isolated nucleic acid molecules are provided encoding novel polypeptides. Novel polypeptides and antibodies that bind to these polypeptides are provided. Also provided are vectors, host cells, and recombinant and synthetic methods for producing human polynucleotides and/or polypeptides, and antibodies. The invention further relates to diagnostic and therapeutic methods useful for diagnosing, treating, preventing and/or prognosing disorders related to these novel polypeptides. The invention further relates to screening methods for identifying agonists and antagonists of polynucleotides and polypeptides of the invention. The present invention further relates to methods and/or compositions for inhibiting or enhancing the production and function of the polypeptides of the present invention.

Detailed Description

Tables

[0041] Table 1 summarizes some of the polynucleotides encompassed by the invention (including cDNA clones related to the sequences (Clone ID NO:Z), contig sequences

(contig identifier (Contig ID:) and contig nucleotide sequence identifier (SEQ ID NO:X)) and further summarizes certain characteristics of these polynucleotides and the polypeptides encoded thereby. The first column provides the gene number in the application for each clone identifier. The second column provides a unique clone identifier, "Clone ID NO:Z", for a cDNA clone related to each contig sequence disclosed in Table 1. The third column provides a unique contig identifier, "Contig ID:" for each of the contig sequences disclosed in Table 1. The fourth column provides the sequence identifier, "SEQ ID NO:X", for each of the contig sequences disclosed in Table 1. The fifth column, "ORF (From-To)", provides the location (i.e., nucleotide position numbers) within the polynucleotide sequence of SEQ ID NO:X that delineate the preferred open reading frame (ORF) that encodes the amino acid sequence shown in the sequence listing and referenced in Table 1 as SEQ ID NO:Y (column 6). Column 7 lists residues comprising predicted epitopes contained in the polypeptides encoded by each of the preferred ORFs (SEQ ID NO:Y). Identification of potential immunogenic regions was performed according to the method of Jameson and Wolf (CABIOS, 4; 181-186 (1988)); specifically, the Genetics Computer Group (GCG) implementation of this algorithm, embodied in the program PEPTIDESTRUCTURE (Wisconsin Package v10.0, Genetics Computer Group (GCG), Madison, Wisc.). This method returns a measure of the probability that a given residue is found on the surface of the protein. Regions where the antigenic index score is greater than 0.9 over at least 6 amino acids are indicated in Table 1 as "Predicted Epitopes". In particular embodiments, polypeptides of the invention comprise, or alternatively consist of, one, two, three, four, five or more of the predicted epitopes described in Table 1. It will be appreciated that depending on the analytical criteria used to predict antigenic determinants, the exact address of the determinant may vary slightly. Column 8, "Tissue Distribution" shows the expression profile of tissue, cells, and/or cell line libraries which express the polynucleotides of the invention. The first number in column 8 (preceding the colon), represents the tissue/cell source identifier code corresponding to the key provided in Table 4. Expression of these polynucleotides was not observed in the other tissues and/or cell libraries tested. For those identifier codes in which the first two letters are not "AR", the second number in column 8 (following the colon), represents the number of times a sequence corresponding to the reference polynucleotide sequence (e.g., SEQ ID NO:X) was identified in the tissue/cell source.

Those tissue/cell source identifier codes in which the first two letters are "AR" designate information generated using DNA array technology. Utilizing this technology, cDNAs were amplified by PCR and then transferred, in duplicate, onto the array. Gene expression was assayed through hybridization of first strand cDNA probes to the DNA array. cDNA probes were generated from total RNA extracted from a variety of different tissues and cell lines. Probe synthesis was performed in the presence of ^{33}P dCTP, using oligo(dT) to prime reverse transcription. After hybridization, high stringency washing conditions were employed to remove non-specific hybrids from the array. The remaining signal, emanating from each gene target, was measured using a Phosphorimager. Gene expression was reported as Phosphor Stimulating Luminescence (PSL) which reflects the level of phosphor signal generated from the probe hybridized to each of the gene targets represented on the array. A local background signal subtraction was performed before the total signal generated from each array was used to normalize gene expression between the different hybridizations. The value presented after "[array code]:" represents the mean of the duplicate values, following background subtraction and probe normalization. One of skill in the art could routinely use this information to identify normal and/or diseased tissue(s) which show a predominant expression pattern of the corresponding polynucleotide of the invention or to identify polynucleotides which show predominant and/or specific tissue and/or cell expression. Column 9 provides the chromosomal location of polynucleotides corresponding to SEQ ID NO:X. Chromosomal location was determined by finding exact matches to EST and cDNA sequences contained in the NCBI (National Center for Biotechnology Information) UniGene database. Given a presumptive chromosomal location, disease locus association was determined by comparison with the Morbid Map, derived from Online Mendelian Inheritance in Man (Online Mendelian Inheritance in Man, OMIMTM, McKusick-Nathans Institute for Genetic Medicine, Johns Hopkins University (Baltimore, MD) and National Center for Biotechnology Information, National Library of Medicine (Bethesda, MD) 2000. World Wide Web URL: <http://www.ncbi.nlm.nih.gov/omim/>). If the putative chromosomal location of the Query overlaps with the chromosomal location of a Morbid Map entry, an OMIM identification number is disclosed in column 10 labeled "OMIM Disease Reference(s)". A key to the OMIM reference identification numbers is provided in Table 5.

[0042] Table 2 summarizes homology and features of some of the polypeptides of the invention. The first column provides a unique clone identifier, "Clone ID NO:Z", corresponding to a cDNA clone disclosed in Table 1. The second column provides the unique contig identifier, "Contig ID:" corresponding to contigs in Table 1 and allowing for correlation with the information in Table 1. The third column provides the sequence identifier, "SEQ ID NO:X", for the contig polynucleotide sequence. The fourth column provides the analysis method by which the homology/identity disclosed in the Table was determined. Comparisons were made between polypeptides encoded by the polynucleotides of the invention and either a non-redundant protein database (herein referred to as "NR"), or a database of protein families (herein referred to as "PFAM") as further described below. The fifth column provides a description of the PFAM/NR hit having a significant match to a polypeptide of the invention. Column six provides the accession number of the PFAM/NR hit disclosed in the fifth column. Column seven, "Score/Percent Identity", provides a quality score or the percent identity, of the hit disclosed in columns five and six. Columns 8 and 9, "NT From" and "NT To" respectively, delineate the polynucleotides in "SEQ ID NO:X" that encode a polypeptide having a significant match to the PFAM/NR database as disclosed in the fifth and sixth columns. In specific embodiments polypeptides of the invention comprise, or alternatively consist of, an amino acid sequence encoded by a polynucleotide in SEQ ID NO:X as delineated in columns 8 and 9, or fragments or variants thereof.

[0043] Table 3 provides polynucleotide sequences that may be disclaimed according to certain embodiments of the invention. The first column provides a unique clone identifier, "Clone ID", for a cDNA clone related to contig sequences disclosed in Table 1. The second column provides the sequence identifier, "SEQ ID NO:X", for contig sequences disclosed in Table 1. The third column provides the unique contig identifier, "Contig ID:", for contigs disclosed in Table 1. The fourth column provides a unique integer 'a' where 'a' is any integer between 1 and the final nucleotide minus 15 of SEQ ID NO:X, and the fifth column provides a unique integer 'b' where 'b' is any integer between 15 and the final nucleotide of SEQ ID NO:X, where both a and b correspond to the positions of nucleotide residues shown in SEQ ID NO:X, and where b is greater than or equal to a + 14. For each of the polynucleotides shown as SEQ ID NO:X, the uniquely defined integers can be substituted into the general formula of a-b, and used to describe polynucleotides

which may be preferably excluded from the invention. In certain embodiments, preferably excluded from the invention are at least one, two, three, four, five, ten, or more of the polynucleotide sequence(s) having the accession number(s) disclosed in the sixth column of this Table (including for example, published sequence in connection with a particular BAC clone). In further embodiments, preferably excluded from the invention are the specific polynucleotide sequence(s) contained in the clones corresponding to at least one, two, three, four, five, ten, or more of the available material having the accession numbers identified in the sixth column of this Table (including for example, the actual sequence contained in an identified BAC clone).

[0044] Table 4 provides a key to the tissue/cell source identifier code disclosed in Table 1, column 8. Column 1 provides the tissue/cell source identifier code disclosed in Table 1, Column 8. Columns 2-5 provide a description of the tissue or cell source. Codes corresponding to diseased tissues are indicated in column 6 with the word "disease". The use of the word "disease" in column 6 is non-limiting. The tissue or cell source may be specific (e.g. a neoplasm), or may be disease-associated (e.g., a tissue sample from a normal portion of a diseased organ). Furthermore, tissues and/or cells lacking the "disease" designation may still be derived from sources directly or indirectly involved in a disease state or disorder, and therefore may have a further utility in that disease state or disorder. In numerous cases where the tissue/cell source is a library, column 7 identifies the vector used to generate the library.

[0045] Table 5 provides a key to the OMIM reference identification numbers disclosed in Table 1, column 10. OMIM reference identification numbers (Column 1) were derived from Online Mendelian Inheritance in Man (Online Mendelian Inheritance in Man, OMIM. McKusick-Nathans Institute for Genetic Medicine, Johns Hopkins University (Baltimore, MD) and National Center for Biotechnology Information, National Library of Medicine, (Bethesda, MD) 2000. World Wide Web URL: <http://www.ncbi.nlm.nih.gov/omim/>). Column 2 provides diseases associated with the cytologic band disclosed in Table 1, column 9, as determined using the Morbid Map database.

Definitions

[0046] The following definitions are provided to facilitate understanding of certain terms used throughout this specification.

[0047] In the present invention, "isolated" refers to material removed from its original environment (e.g., the natural environment if it is naturally occurring), and thus is altered "by the hand of man" from its natural state. For example, an isolated polynucleotide could be part of a vector or a composition of matter, or could be contained within a cell, and still be "isolated" because that vector, composition of matter, or particular cell is not the original environment of the polynucleotide. The term "isolated" does not refer to genomic or cDNA libraries, whole cell total or mRNA preparations, genomic DNA preparations (including those separated by electrophoresis and transferred onto blots), sheared whole cell genomic DNA preparations or other compositions where the art demonstrates no distinguishing features of the polynucleotide/sequences of the present invention.

[0048] As used herein, a "polynucleotide" refers to a molecule having a nucleic acid sequence encoding SEQ ID NO:Y or a fragment or variant thereof; a nucleic acid sequence contained in SEQ ID NO:X (as described in column 3 of Table 1) or the complement thereof; or a cDNA sequence contained in Clone ID NO:Z (as described in column 2 of Table 1 and contained within ATCC Deposit No. PTA-2448 or PTA-2449). For example, the polynucleotide can contain the nucleotide sequence of the full length cDNA sequence, including the 5' and 3' untranslated sequences, the coding region, as well as fragments, epitopes, domains, and variants of the nucleic acid sequence. Moreover, as used herein, a "polypeptide" refers to a molecule having an amino acid sequence encoded by a polynucleotide of the invention as broadly defined (obviously excluding poly-Phenylalanine or poly-Lysine peptide sequences which result from translation of a polyA tail of a sequence corresponding to a cDNA).

[0049] In the present invention, "SEQ ID NO:X" was often generated by overlapping sequences contained in multiple clones (contig analysis). A representative clone containing all or most of the sequence for SEQ ID NO:X is deposited at Human Genome Sciences, Inc. (HGS) in a catalogued and archived library. As shown, for example, in column 2 of Table 1, each clone is identified by a cDNA Clone ID (identifier generally referred to herein as Clone ID NO:Z). Each Clone ID is unique to an individual clone and the Clone ID is all the information needed to retrieve a given clone from the HGS library.

Furthermore, certain clones disclosed in this application have been deposited with the ATCC on September 12, 2000 and give ATCC Designation Numbers PTA-2448 and PTA-2449. The ATCC is located at 10801 University Boulevard, Manassas, Virginia 20110-2209, USA. The ATCC deposits were made pursuant to the terms of the Budapest Treaty on the international recognition of the deposit of microorganisms for the purposes of patent procedure.

[0050] In specific embodiments, the polynucleotides of the invention are at least 15, at least 30, at least 50, at least 100, at least 125, at least 500, or at least 1000 continuous nucleotides but are less than or equal to 300 kb, 200 kb, 100 kb, 50 kb, 15 kb, 10 kb, 7.5kb, 5 kb, 2.5 kb, 2.0 kb, or 1 kb, in length. In a further embodiment, polynucleotides of the invention comprise a portion of the coding sequences, as disclosed herein, but do not comprise all or a portion of any intron. In another embodiment, the polynucleotides comprising coding sequences do not contain coding sequences of a genomic flanking gene (i.e., 5' or 3' to the gene of interest in the genome). In other embodiments, the polynucleotides of the invention do not contain the coding sequence of more than 1000, 500, 250, 100, 50, 25, 20, 15, 10, 5, 4, 3, 2, or 1 genomic flanking gene(s).

[0051] A "polynucleotide" of the present invention also includes those polynucleotides capable of hybridizing, under stringent hybridization conditions, to sequences contained in SEQ ID NO:X, or the complement thereof (e.g., the complement of any one, two, three, four, or more of the polynucleotide fragments described herein), the polynucleotide sequence delineated in columns 8 and 9 of Table 2 or the complement thereof, and/or cDNA sequences contained in Clone ID NO:Z (e.g., the complement of any one, two, three, four, or more of the polynucleotide fragments, or the cDNA clone within the pool of cDNA clones deposited with the ATCC, described herein). "Stringent hybridization conditions" refers to an overnight incubation at 42 degree C in a solution comprising 50% formamide, 5x SSC (750 mM NaCl, 75 mM trisodium citrate), 50 mM sodium phosphate (pH 7.6), 5x Denhardt's solution, 10% dextran sulfate, and 20 μ g/ml denatured, sheared salmon sperm DNA, followed by washing the filters in 0.1x SSC at about 65 degree C.

[0052] Also contemplated are nucleic acid molecules that hybridize to the polynucleotides of the present invention at lower stringency hybridization conditions. Changes in the stringency of hybridization and signal detection are primarily accomplished through the manipulation of formamide concentration (lower percentages of

formamide result in lowered stringency); salt conditions, or temperature. For example, lower stringency conditions include an overnight incubation at 37 degree C in a solution comprising 6X SSPE (20X SSPE = 3M NaCl; 0.2M NaH₂PO₄; 0.02M EDTA, pH 7.4), 0.5% SDS, 30% formamide, 100 ug/ml salmon sperm blocking DNA; followed by washes at 50 degree C with 1XSSPE, 0.1% SDS. In addition, to achieve even lower stringency, washes performed following stringent hybridization can be done at higher salt concentrations (e.g. 5X SSC).

[0053] Note that variations in the above conditions may be accomplished through the inclusion and/or substitution of alternate blocking reagents used to suppress background in hybridization experiments. Typical blocking reagents include Denhardt's reagent, BLOTTO, heparin, denatured salmon sperm DNA, and commercially available proprietary formulations. The inclusion of specific blocking reagents may require modification of the hybridization conditions described above, due to problems with compatibility.

[0054] Of course, a polynucleotide which hybridizes only to polyA+ sequences (such as any 3' terminal polyA+ tract of a cDNA shown in the sequence listing), or to a complementary stretch of T (or U) residues, would not be included in the definition of "polynucleotide," since such a polynucleotide would hybridize to any nucleic acid molecule containing a poly (A) stretch or the complement thereof (e.g., practically any double-stranded cDNA clone generated using oligo dT as a primer).

[0055] The polynucleotide of the present invention can be composed of any polyribonucleotide or polydeoxribonucleotide, which may be unmodified RNA or DNA or modified RNA or DNA. For example, polynucleotides can be composed of single- and double-stranded DNA, DNA that is a mixture of single- and double-stranded regions, single- and double-stranded RNA, and RNA that is mixture of single- and double-stranded regions, hybrid molecules comprising DNA and RNA that may be single-stranded or, more typically, double-stranded or a mixture of single- and double-stranded regions. In addition, the polynucleotide can be composed of triple-stranded regions comprising RNA or DNA or both RNA and DNA. A polynucleotide may also contain one or more modified bases or DNA or RNA backbones modified for stability or for other reasons. "Modified" bases include, for example, tritylated bases and unusual bases such as inosine.

A variety of modifications can be made to DNA and RNA; thus, "polynucleotide" embraces chemically, enzymatically, or metabolically modified forms.

[0056] The polypeptide of the present invention can be composed of amino acids joined to each other by peptide bonds or modified peptide bonds, i.e., peptide isosteres, and may contain amino acids other than the 20 gene-encoded amino acids. The polypeptides may be modified by either natural processes, such as posttranslational processing, or by chemical modification techniques which are well known in the art. Such modifications are well described in basic texts and in more detailed monographs, as well as in a voluminous research literature. Modifications can occur anywhere in a polypeptide, including the peptide backbone, the amino acid side-chains and the amino or carboxyl termini. It will be appreciated that the same type of modification may be present in the same or varying degrees at several sites in a given polypeptide. Also, a given polypeptide may contain many types of modifications. Polypeptides may be branched, for example, as a result of ubiquitination, and they may be cyclic, with or without branching. Cyclic, branched, and branched cyclic polypeptides may result from posttranslation natural processes or may be made by synthetic methods. Modifications include acetylation, acylation, ADP-ribosylation, amidation, covalent attachment of flavin, covalent attachment of a heme moiety, covalent attachment of a nucleotide or nucleotide derivative, covalent attachment of a lipid or lipid derivative, covalent attachment of phosphatidylinositol, cross-linking, cyclization, disulfide bond formation, demethylation, formation of covalent cross-links, formation of cysteine, formation of pyroglutamate, formylation, gamma-carboxylation, glycosylation, GPI anchor formation, hydroxylation, iodination, methylation, myristoylation, oxidation, pegylation, proteolytic processing, phosphorylation, prenylation, racemization, selenoylation, sulfation, transfer-RNA mediated addition of amino acids to proteins such as arginylation, and ubiquitination. (See, for instance, PROTEINS - STRUCTURE AND MOLECULAR PROPERTIES, 2nd Ed., T. E. Creighton, W. H. Freeman and Company, New York (1993); POSTTRANSLATIONAL COVALENT MODIFICATION OF PROTEINS, B. C. Johnson, Ed., Academic Press, New York, pgs. 1-12 (1983); Seifter et al., Meth. Enzymol. 182:626-646 (1990); Rattan et al., Ann. N.Y. Acad. Sci. 663:48-62 (1992)).

[0057] "SEQ ID NO:X" refers to a polynucleotide sequence described, for example, in Tables 1 or 2, while "SEQ ID NO:Y" refers to a polypeptide sequence described in

column 6 of Table 1. SEQ ID NO:X is identified by an integer specified in column 4 of Table 1. The polypeptide sequence SEQ ID NO:Y is a translated open reading frame (ORF) encoded by polynucleotide SEQ ID NO:X. "Clone ID NO:Z" refers to a cDNA clone described in column 2 of Table 1.

[0058] "A polypeptide having functional activity" refers to a polypeptide capable of displaying one or more known functional activities associated with a full-length (complete) protein. Such functional activities include, but are not limited to, biological activity, antigenicity [ability to bind (or compete with a polypeptide for binding) to an anti-polypeptide antibody], immunogenicity (ability to generate antibody which binds to a specific polypeptide of the invention), ability to form multimers with polypeptides of the invention, and ability to bind to a receptor or ligand for a polypeptide.

[0059] The polypeptides of the invention can be assayed for functional activity (e.g. biological activity) using or routinely modifying assays known in the art, as well as assays described herein. Specifically, one of skill in the art may routinely assay polypeptides (including fragments and variants) of the invention for activity using assays as described in the Examples

[0060] "A polypeptide having biological activity" refers to a polypeptide exhibiting activity similar to, but not necessarily identical to, an activity of a polypeptide of the present invention, including mature forms, as measured in a particular biological assay, with or without dose dependency. In the case where dose dependency does exist, it need not be identical to that of the polypeptide, but rather substantially similar to the dose-dependence in a given activity as compared to the polypeptide of the present invention (i.e., the candidate polypeptide will exhibit greater activity or not more than about 25-fold less and, preferably, not more than about tenfold less activity, and most preferably, not more than about three-fold less activity relative to the polypeptide of the present invention).

[0061] Table 1 summarizes some of the polynucleotides encompassed by the invention (including contig sequences (SEQ ID NO:X) and clones (Clone ID NO:Z) and further summarizes certain characteristics of these polynucleotides and the polypeptides encoded thereby.

Polynucleotides and Polypeptides of the InventionTABLE 1

Gene No:	Clone ID NO: Z	Contig ID:	SEQ ID NO: X	ORF (From-To)	AA SEQ ID NO: Y	Predicted Epitopes	Tissue Distribution Library code: count (see Table IV for Library Codes)	Cytologic Band	OMIM Disease Reference(s):
1	HEMEC34	684296	11	55 - 879	187	Ala-18 to Ser-24, Glu-44 to Ala-57, Ala-114 to Arg-120, Thr-129 to Gln-137, Thr-152 to Asn-160, Ser-187 to Asp-199, Thr-209 to Pro-215.	AR241: 16, AR313: 13, AR192: 9, AR312: 8, AR219: 8, AR218: 8, AR204: 7, AR271: 7, AR251: 7, AR182: 7, AR282: 7, AR229: 7, AR198: 7, AR184: 6, AR263: 6, AR096: 6, AR269: 6, AR293: 6, AR089: 6, AR060: 6, AR258: 6, AR296: 6, AR183: 6, AR213: 6, AR185: 6, AR053: 5, AR299: 5, AR240: 5, AR243: 5, AR273: 5, AR039: 5, AR238: 5, AR300: 5, AR206: 5, AR226: 5, AR316: 5, AR290: 5, AR052: 5,		

									AR275: 4, AR286: 4, AR246: 4, AR186: 4, AR104: 4, AR247: 4, AR270: 4, AR310: 4, AR205: 4, AR267: 4, AR194: 4, AR285: 4, AR177: 4, AR268: 4, AR266: 4, AR291: 4, AR233: 4, AR298: 4, AR274: 4, AR175: 4, AR055: 4, AR202: 3, AR033: 3, AR294: 3, AR292: 3, AR259: 3, AR237: 3, AR289: 3, AR277: 3, AR295: 3, AR234: 3, AR061: 2, AR227: 2, AR284: 2, AR253: 2, AR309: 2, AR231: 2, AR256: 2, AR283: 2, AR179: 2, AR232: 1, AR314: 1, AR315: 1, AR249: 1, AR265: 1, AR248: 1, AR280: 1, AR244: 0, AR281: 0 S0046: 4, H0623: 3, S0045: 1, H0069: 1, H0457: 1, H0109: 1, S0036: 1, H0634: 1, H0056: 1 and S0152: 1.
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2	HDPMQ69	812744	12	1 - 669	188	<p>Arg-1 to Ala-9, Pro-11 to Glu-17, Leu-19 to Gly-30, Pro-84 to Gly-90, Asn-114 to Val-120, Ser-123 to Phe-128, Gly-132 to Asn-138, Gln-149 to Tyr-154, Arg-203 to Pro-223.</p>	<p>AR313: 32, AR241: 19, AR248: 19, AR096: 18, AR253: 17, AR312: 14, AR089: 14, AR192: 13, AR104: 13, AR213: 12, AR183: 12, AR300: 12, AR299: 12, AR275: 11, AR247: 11, AR198: 10, AR052: 10, AR218: 10, AR238: 10, AR229: 10, AR249: 10, AR293: 10, AR053: 10, AR271: 10, AR240: 10, AR292: 9, AR274: 9, AR263: 8, AR185: 8, AR204: 8, AR316: 8, AR243: 8, AR314: 8, AR258: 8, AR226: 8, AR265: 8, AR219: 7, AR186: 7, AR280: 7, AR251: 7, AR315: 7, AR039: 7, AR033: 7, AR273: 7, AR282: 6, AR060: 6, AR309: 6, AR175: 6, AR277: 6, AR294: 6, AR244: 6, AR177: 6, AR234: 6, AR194: 5, AR295: 5, AR310: 5, AR231: 5, AR206: 5, AR233: 5, AR184: 5,</p>		
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3	HTPAU91	846654	13	1282 - 2262	189	Thr-42 to Ser-50, Gly-88 to Pro-97, His-106 to Asp-118, Arg-198 to Ser-214, Asn-225 to Asn-231, Ser-291 to Cys-297, Asn-319 to Asn-324.	AR248: 10, AR205: 7, AR215: 6, AR235: 5, AR298: 4, AR249: 4, AR253: 3, AR251: 3, AR247: 3, AR170: 3, AR171: 3, AR282: 2, AR225: 2, AR245: 2, AR089: 2, AR312: 2, AR224: 2, AR264: 2, AR096: 2, AR240: 2, AR295: 2, AR268: 2, AR221: 2, AR299: 1,		

									AR163: 1, AR166: 1, AR296: 1, AR266: 1, AR285: 1, AR313: 1, AR217: 1, AR201: 1, AR277: 1, AR257: 1, AR274: 1, AR237: 1, AR309: 1, AR286: 1, AR308: 1, AR283: 1, AR294: 1, AR161: 1, AR162: 1, AR232: 1, AR199: 1, AR175: 1, AR261: 1, AR272: 1, AR289: 1, AR194: 1, AR267: 1, AR181: 1, AR316: 1, AR230: 1, AR300: 1, AR297: 1, AR039: 1, AR061: 1, AR168: 1, AR179: 1, AR202: 1, AR270: 1, AR053: 1, AR236: 1, AR184: 1, AR310: 1, AR182: 1, AR104: 1, AR033: 1, AR213: 1, AR227: 1, AR288: 1, AR060: 1, AR186: 1, AR269: 1, AR180: 1, AR252: 1, AR311: 1, AR210: 1, AR280: 0, AR234: 0, AR185: 0, AR204: 0, AR229: 0.
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L0526: 5, H0658: 5, S0380: 5, H0134: 5, L0740: 5, L0747: 5, L0605: 5, L0362: 5, H0422: 5, S0424: 5, H0686: 4, S0342: 4, S0376: 4, H0351: 4, H0586: 4, L0586: 4, H0250: 4, H0421: 4, H0009: 4, L0163: 4, H0083: 4, H0266: 4, S0022: 4, H0617: 4, H0135: 4, H0087: 4, H0116: 4, H0100: 4, L0662: 4, L0497: 4, L0655: 4, L0527: 4, L0657: 4, L0666: 4, H0666: 4, S0044: 4, H0576: 4, S3012: 4, S3014: 4, L0756: 4, H0171: 3, S6024: 3, H0484: 3, H0669: 3, S0410: 3, H0497: 3, L0623: 3, H0599: 3, H0575: 3, T0048: 3, H0581: 3, H0597: 3, H0039: 3, H0644: 3, H0163: 3, H0551: 3, T0067: 3, T0041: 3, S0422: 3, H0529: 3,
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T0006: 2, S0036: 2,
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H0059: 2, H0625: 2,
H0509: 2, H0646: 2,
S0210: 2, L0598: 2,
L0763: 2, L0500: 2,
L0501: 2, L0549: 2,
L0803: 2, L0376: 2,
L0653: 2, L0542: 2,
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H0187: 2, S0392: 2,
S0390: 2, L0742: 2,
H0444: 2, L0596: 2,
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L0599: 2, L0603: 2,
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H0661: 1, H0663: 1,
H0664: 1, H0589: 1,
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H0489: 1, S0278: 1,
H0369: 1, H0441: 1,
H0632: 1, H0492: 1,
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4	H7TBE17	858079	14	706 - 2	190	Gln-155 to Ser-164.	L0774: 1, L0775: 1, L0651: 1, L0378: 1, L0805: 1, L0509: 1, L0652: 1, L0555: 1, L0513: 1, L0656: 1, L0559: 1, L0384: 1, L0809: 1, L0529: 1, L0368: 1, L0532: 1, H0691: 1, H0520: 1, H0519: 1, H0684: 1, H0651: 1, S0454: 1, S0013: 1, S0406: 1, H0478: 1, L0744: 1, L0439: 1, L0731: 1, H0343: 1, L0592: 1, L0485: 1, L0581: 1, L0608: 1, L0593: 1, H0217: 1, H0216: 1, H0008: 1 and H0352: 1. AR313: 15, AR291: 9, AR249: 9, AR296: 8, AR183: 8, AR312: 8, AR096: 7, AR241: 7, AR270: 7, AR292: 7, AR269: 7, AR229: 6, AR089: 6, AR290: 6, AR258: 6, AR268: 6, AR184: 6, AR293: 6, AR219: 6, AR182: 6, AR298: 5, AR247: 5,		
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5	H7TRA52	858086	15	703 - 2	191	Gln-154 to Ser-163.	AR274: 7, AR312: 6, AR273: 6, AR310: 6, AR053: 6, AR309: 6, AR251: 6, AR313: 5, AR271: 5, AR241: 4, AR263: 4, AR213: 4, AR052: 4, AR243: 4, AR265: 4, AR247: 3, AR282: 3, AR089: 3, AR096: 3, AR299: 3, AR183: 3, AR248: 3, AR175: 3, AR218: 3, AR219: 2, AR185: 2, AR275: 2, AR060: 2, AR292: 2, AR039: 2, AR240: 2, AR280: 2, AR256: 2, AR316: 2, AR277: 2, AR186: 2, AR033: 2, AR315: 2, AR104: 2, AR283: 2, AR253: 2, AR205: 2, AR314: 2, AR229: 2, AR177: 2, AR249: 2,				

6	HSSJC65	872582	16	3 - 1688	192	His-1 to Ala-16, Thr-53 to Gly-59, Pro-182 to Asp-188, Leu-269 to Trp-274, Gln-288 to Ser-295,	AR293: 2, AR198: 2, AR300: 1, AR204: 1, AR295: 1, AR055: 1, AR290: 1, AR192: 1, AR238: 1, AR291: 1, AR237: 1, AR179: 1, AR061: 1, AR232: 1, AR231: 1, AR281: 1, AR226: 1, AR294: 1, AR296: 1, AR234: 1, AR289: 1, AR268: 1, AR227: 1, AR244: 1, AR258: 1, AR270: 1, AR259: 1, AR298: 1, AR284: 1, AR286: 1, AR202: 1, AR269: 1, AR266: 1, AR233: 0, AR267: 0, AR194: 0, AR285: 0, AR246: 0, AR206: 0, AR182: 0, AR184: 0 S0198: 54, S0274: 12, S0252: 4, S0270: 3, S0264: 1, S0268: 1 and S0228: 1.	22q11	104170, 104170, 104170, 115470, 142360,
							AR298: 5, AR033: 5, AR293: 4, AR104: 4, AR244: 4, AR192: 4, AR186: 3, AR231: 3, AR182: 3, AR061: 3,		

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7	HKLAA30	875001	17	64 - 222	193		AR253: 0, AR283: 0, AR241: 0, AR271: 0, AR281: 0, AR219: 0, AR273: 0, AR309: 0, AR039: 0, AR226: 0, AR280: 0, AR204: 0, AR248: 0 H0052: 3, L0439: 3, H0194: 1, H0546: 1, H0615: 1, H0135: 1, H0087: 1, L0759: 1 and S0456: 1.	2p25	274500, 274500, 274500, 602134
							AR271: 157, AR196: 101, AR191: 101, AR274: 98, AR211: 87, AR190: 83, AR244: 81, AR245: 77, AR189: 77, AR174: 76, AR188: 76, AR060: 75, AR261: 70, AR235: 66, AR177: 64, AR173: 63, AR175: 61, AR161: 60, AR162: 60, AR275: 60, AR254: 59, AR264: 58, AR163: 58, AR165: 58, AR096: 56, AR197: 56, AR192: 56, AR246: 56, AR164: 55, AR250: 55, AR240: 54, AR210: 54, AR236: 51, AR195: 51, AR178: 51,		

	AR104: 50, AR311: 50, AR206: 49, AR180: 49, AR166: 49, AR182: 48, AR272: 48, AR176: 48, AR181: 47, AR243: 47, AR290: 47, AR270: 47, AR297: 47, AR186: 46, AR295: 46, AR273: 46, AR287: 46, AR312: 45, AR269: 45, AR089: 44, AR309: 44, AR248: 44, AR268: 43, AR198: 43, AR039: 43, AR033: 43, AR263: 42, AR288: 41, AR291: 40, AR184: 39, AR262: 39, AR194: 39, AR249: 39, AR207: 39, AR266: 39, AR252: 38, AR253: 38, AR183: 38, AR316: 38, AR256: 38, AR200: 37, AR247: 37, AR265: 36, AR251: 36, AR255: 36, AR218: 35, AR313: 35, AR285: 35, AR219: 35, AR308: 34, AR239: 34, AR201: 34, AR282: 33, AR310: 33, AR193: 33, AR296: 33, AR185: 32, AR299: 32, AR052: 32, AR260: 32,	
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[illegible]

8	HPICP59	943763	18	202 - 1083	194	Asp-1 to Met-7, Leu-62 to Tyr-68, Glu-79 to Asn-87.	L0777: 1. AR273: 8, AR241: 7, AR052: 7, AR312: 6, AR313: 6, AR274: 5, AR271: 5, AR309: 5, AR219: 5, AR275: 4, AR218: 4, AR213: 4, AR186: 4, AR310: 4, AR053: 4, AR039: 4, AR248: 4, AR299: 4, AR060: 4, AR096: 3, AR183: 3, AR089: 3, AR269: 3, AR277: 3, AR267: 3, AR175: 3, AR253: 3, AR265: 3, AR198: 3, AR263: 3, AR240: 3, AR185: 3, AR205: 3, AR316: 3, AR268: 3, AR293: 3, AR300: 3, AR229: 3, AR266: 2, AR292: 2, AR182: 2, AR206: 2, AR282: 2, AR184: 2, AR246: 2, AR177: 2, AR231: 2, AR249: 2, AR270: 2, AR280: 2, AR055: 2, AR104: 2, AR033: 2, AR247: 2, AR238: 2, AR243: 2, AR061: 2, AR295: 2,		
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9	HTEBO72	947922	19	27 - 575	195	Arg-32 to Glu-43, Met-144 to Asn-150, His-160 to Lys-169.	AR232: 2, AR237: 2, AR290: 2, AR289: 2, AR234: 2, AR294: 2, AR226: 1, AR283: 1, AR194: 1, AR179: 1, AR233: 1, AR258: 1, AR286: 1, AR314: 1, AR284: 1, AR259: 1, AR298: 1, AR285: 1, AR192: 1, AR296: 1, AR315: 1, AR251: 1, AR256: 1, AR244: 1, AR227: 0, AR204: 0, AR202: 0, AR291: 0, AR281: 0 H0328: 1, S0208: 1, S0152: 1 and H0521: 1.	8p22	148370, 238600, 238600, 238600, 238600, 600143, 601385, 602629
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[illegible]

10	HWLHQ59	949724	20	1205 - 282	196	Arg-1 to Asp-7, Glu-22 to Ser-31, Leu-47 to Gly-61.	<p>135, AR218: 130, AR183: 128, AR053: 119, AR269: 114, AR096: 98, AR249: 94, AR266: 93, AR052: 93, AR310: 93, AR277: 90, AR268: 88, AR289: 85, AR309: 83, AR213: 83, AR270: 80, AR251: 78, AR253: 71, AR248: 67, AR312: 67, AR265: 66, AR263: 58, AR313: 58 L0779: 4, H0038: 3 and L0758: 3.</p> <p>AR281: 27, AR280: 18, AR315: 18, AR314: 15, AR263: 13, AR265: 11, AR310: 11, AR283: 11, AR247: 9, AR033: 8, AR251: 8, AR300: 7, AR299: 7, AR295: 7, AR096: 7, AR282: 7, AR277: 6, AR089: 6, AR232: 6, AR312: 5, AR183: 5, AR313: 5, AR229: 5, AR248: 5, AR316: 5, AR285: 5, AR177: 5, AR266: 5, AR055: 5, AR267: 5, AR238: 5, AR234: 4,</p>		
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AR253:	4, AR309:	4,
AR227:	4, AR194:	4,
AR291:	4, AR231:	4,
AR226:	4, AR292:	4,
AR294:	4, AR289:	4,
AR269:	4, AR237:	4,
AR284:	4, AR286:	4,
AR296:	4, AR213:	4,
AR298:	4, AR175:	3,
AR233:	3, AR293:	3,
AR249:	3, AR052:	3,
AR270:	3, AR182:	3,
AR268:	3, AR218:	3,
AR258:	3, AR219:	3,
AR061:	3, AR243:	3,
AR256:	3, AR184:	3,
AR053:	2, AR060:	2,
AR290:	2, AR202:	2,
AR259:	2, AR246:	2,
AR185:	2, AR198:	2,
AR275:	2, AR179:	2,
AR206:	2, AR271:	1,
AR186:	1, AR104:	1,
AR240:	1, AR274:	1,
AR204:	0, AR039:	0,
AR273:	0, AR192:	0,
AR244:	0, AR205:	0,
AR241:	0	
	S0358: 6, S0354: 5,	
	S0374: 2 and H0675: 1.	

11	HDQES11	950838	21	2042 - 282	197	<p>Leu-12 to Arg-21, Pro-23 to Pro-28, Asp-85 to Arg-90, Gln-147 to Glu-152, Lys-176 to Gly-185, Glu-230 to Leu-237, Leu-255 to His-262, Lys-276 to His-284, Ser-292 to Lys-303, Pro-342 to Tyr-351, Asn-393 to Arg-399, Gly-442 to Gly-447, Thr-478 to Met-485, Pro-488 to Tyr-495, Glu-505 to Asn-512.</p>	<p>AR309: 46, AR052: 42, AR265: 41, AR312: 36, AR313: 34, AR263: 33, AR310: 33, AR213: 31, AR281: 29, AR194: 29, AR274: 28, AR280: 27, AR053: 26, AR314: 25, AR271: 25, AR241: 24, AR205: 24, AR315: 24, AR266: 24, AR273: 23, AR270: 23, AR268: 22, AR269: 22, AR039: 22, AR246: 22, AR286: 21, AR275: 20, AR285: 20, AR096: 20, AR198: 20, AR192: 20, AR202: 19, AR296: 19, AR206: 19, AR104: 18, AR089: 18, AR291: 17, AR033: 17, AR243: 17, AR248: 17, AR253: 17, AR290: 17, AR316: 17, AR184: 17, AR289: 16, AR247: 16, AR244: 16, AR284: 16, AR299: 16, AR182: 15, AR298: 15, AR267: 15, AR282: 15, AR283: 14, AR249: 13, AR204: 13, AR300: 13, AR234: 12, AR186: 12, AR060: 12,</p>	15q22	102578, 109700, 151670, 154550, 601780
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							AR240: 12, AR231: 12, AR238: 12, AR251: 11, AR277: 11, AR183: 11, AR232: 11, AR229: 10, AR055: 10, AR185: 9, AR258: 8, AR175: 8, AR226: 8, AR227: 8, AR294: 8, AR219: 8, AR293: 7, AR061: 7, AR292: 7, AR218: 7, AR237: 7, AR177: 6, AR259: 6, AR295: 5, AR233: 5, AR256: 4, AR179: 2 L0766: 8, H0521: 4, S0358: 3, H0543: 3, T0042: 2, L0521: 2, L0517: 2, S0114: 1, S0282: 1, S6016: 1, S0280: 1, H0545: 1, H0123: 1, L0456: 1, H0090: 1, H0616: 1, L0646: 1, L0775: 1, L0664: 1, H0547: 1, L0750: 1, L0779: 1, L0601: 1 and S0276: 1.		
12	HOHCE96	955622	22	2 - 904	198	Ser-4 to Lys-13, Ser-26 to Tyr-33, Asp-104 to Gln-112, Pro-168 to Pro-174,	AR180: 6, AR176: 3, AR215: 3, AR171: 2, AR163: 2, AR221: 2, AR263: 2, AR192: 2,		

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AR254:	0, AR055:	0,						
AR177:	0, AR238:	0,						
AR271:	0, AR181:	0,						
AR195:	0, AR247:	0,						
AR189:	0, AR249:	0,						
AR256:	0, AR241:	0,						
AR264:	0, AR173:	0,						
AR039:	0, AR265:	0,						
AR259:	0, AR194:	0,						
AR289:	0, AR218:	0,						
AR260:	0, AR255:	0,						
AR202:	0, AR300:	0,						
AR252:	0, AR314:	0,						
AR052:	0, AR246:	0,						
AR211:	0, AR296:	0,						
AR205:	0, AR258:	0,						
AR053:	0, AR207:	0,						
AR269:	0, AR198:	0,						
AR298:	0, AR206:	0,						
AR297:	0, AR193:	0,						
AR286:	0, AR061:	0,						
AR251:	0, AR213:	0,						
AR201:	0, AR243:	0,						
AR293:	0, AR233:	0,						
AR203:	0, AR190:	0,						
AR219:	0, AR169:	0,						
AR191:	0, AR229:	0,						
AR234:	0, AR222:	0,						
AR284:	0, AR199:	0,						
AR237:	0, AR261:	0,						

13	HBKDR69	956894	23	1044 - 490	199			AR285: 0, AR231: 0, AR242: 0, AR236: 0, AR230: 0, AR184: 0, AR239: 0 H0030: 2, S0250: 1, L0565: 1 and S0276: 1.	5q31		121050, 131400, 138040, 153455, 159000, 179095, 181460, 192974, 192974, 600807, 601596, 601692, 601692, 601692, 601692, 602089, 602121, 602460
								Lys-41 to Lys-46, Phe-68 to Glu-74, Val-100 to Gly-106, Ala-129 to Gln-140, Tyr-180 to Leu-185.			
								AR258: 77, AR292: 62, AR259: 56, AR256: 46, AR294: 45, AR295: 43, AR293: 36, AR283: 27, AR286: 22, AR266: 20, AR291: 17, AR296: 14, AR289: 14, AR298: 13, AR285: 11, AR284: 11, AR316: 5, AR053: 3, AR183: 3, AR290: 3, AR282: 3, AR244: 3, AR206: 2, AR204: 2, AR089: 2, AR310: 2, AR192: 2, AR205: 2, AR313: 2, AR312: 2, AR096: 2, AR104: 2, AR299: 2, AR270: 1, AR060: 1, AR182: 1, AR033: 1, AR052: 1, AR232: 1, AR231: 1, AR186: 1, AR268: 1, AR300: 1, AR241: 1, AR277: 1, AR061: 1, AR237: 1, AR194: 1,			

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14	HSIGI60	956986	24	1251 - 580	200	Asp-28 to Gly-33, Pro-123 to Thr-129, Val-189 to Lys-194, Gln-218 to Glu-224.			AR186: 12, AR194: 10, AR206: 9, AR246: 6, AR202: 6, AR249: 6, AR225: 5, AR244: 5, AR265: 5, AR251: 5, AR180: 5, AR243: 5, AR033: 4, AR310: 4, AR184: 4, AR055: 4,				

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AR229:	1, AR237:	1,
AR178:	1, AR298:	1,
AR274:	1, AR191:	1,
AR177:	1, AR261:	1,
AR296:	1, AR297:	1,
AR284:	1, AR232:	1,
AR222:	1, AR290:	1,
AR182:	1, AR226:	1,
AR227:	1, AR234:	1,
AR236:	1, AR291:	1,
AR258:	1, AR215:	1,
AR286:	1, AR200:	1,
AR280:	1, AR235:	1,
AR219:	1, AR314:	1,
AR231:	1, AR308:	1,
AR256:	1, AR211:	1,
AR168:	1, AR216:	0,
AR263:	0, AR281:	0,
AR218:	0, AR217:	0,
AR207:	0, AR245:	0,
AR266:	0, AR250:	0,
AR176:	0, AR193:	0,
AR242:	0, AR287:	0,
AR179:	0, AR197:	0,
AR264:	0, AR252:	0,
AR315:	0, AR196:	0,
AR181:	0, AR221:	0,
AR188:	0, AR272:	0,
AR210:	0, AR173:	0,
AR190:	0, AR199:	0,

15	HOFNI12	982193	25	3 - 362	201	Asp-1 to Asp-7, Leu-12 to Ser-21, Thr-41 to Gly-46.	AR260: 0, AR195: 0, AR255: 0, AR162: 0, AR203: 0, AR262: 0 L0439: 5, L0756: 5, H0590: 4, L0438: 2, S0046: 1 and L0665: 1. AR292: 6, AR183: 4, AR293: 4, AR241: 4, AR269: 3, AR039: 3, AR182: 3, AR296: 3, AR256: 3, AR253: 3, AR184: 3, AR274: 3, AR266: 3, AR270: 3, AR089: 2, AR282: 2, AR271: 2, AR294: 2, AR295: 2, AR258: 2, AR291: 2, AR186: 2, AR096: 2, AR286: 2, AR313: 2, AR300: 2, AR298: 2, AR299: 2, AR259: 2, AR175: 2, AR309: 2, AR052: 2, AR179: 1, AR268: 1, AR289: 1, AR284: 1, AR316: 1, AR247: 1, AR060: 1, AR240: 1, AR290: 1, AR285: 1, AR185: 1, AR234: 1, AR244: 1, AR219: 1, AR033: 1, AR177: 1,			
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16	HEGAZ61	983192	26	12 - 560	202	Leu-45 to Lys-50, Gln-55 to Glu-74.	AR104: 1, AR283: 1, AR227: 1, AR061: 1, AR275: 1, AR218: 1, AR226: 1, AR312: 1, AR192: 1, AR277: 1, AR314: 1, AR273: 0, AR229: 0, AR281: 0, AR267: 0, AR237: 0, AR204: 0, AR243: 0, AR198: 0, AR280: 0, AR233: 0, AR310: 0, AR202: 0, AR238: 0, AR315: 0, AR263: 0, AR249: 0, AR194: 0, AR251: 0, AR232: 0, AR213: 0, AR231: 0, AR055: 0 H0415: 1		
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AR247:	9, AR316:	9,
AR215:	9, AR230:	9,
AR300:	9, AR104:	9,
AR294:	9, AR210:	9,
AR292:	9, AR283:	9,
AR216:	9, AR226:	9,
AR291:	9, AR262:	9,
AR238:	9, AR298:	9,
AR189:	9, AR296:	8,
AR270:	8, AR240:	8,
AR313:	8, AR182:	8,
AR255:	8, AR089:	8,
AR200:	8, AR175:	8,
AR185:	8, AR229:	8,
AR268:	8, AR237:	8,
AR297:	8, AR293:	7,
AR176:	7, AR252:	7,
AR231:	7, AR248:	7,
AR245:	7, AR269:	7,
AR173:	7, AR258:	7,
AR197:	7, AR272:	7,
AR186:	7, AR193:	7,
AR290:	6, AR249:	6,
AR061:	6, AR256:	6,
AR060:	6, AR234:	6,
AR253:	6, AR218:	6,
AR211:	6, AR250:	6,
AR183:	6, AR179:	6,
AR184:	6, AR267:	6,
AR181:	6, AR259:	6,

17	HHERQ79	985266	27	340 - 726	203	Ser-3 to Thr-11, Lys-32 to Gly-39, Thr-50 to Glu-57, Thr-83 to Gln-88.	AR201: 5, AR219: 5, AR260: 5, AR188: 3, AR178: 2, AR180: 1, AR254: 1 H0549: 1 and H0550: 1. AR265: 7, AR310: 7, AR263: 6, AR268: 5, AR213: 5, AR312: 5, AR309: 4, AR315: 4, AR251: 4, AR281: 4, AR270: 4, AR282: 3, AR267: 3, AR283: 3, AR053: 3, AR299: 3, AR249: 3, AR266: 3, AR183: 3, AR277: 3, AR280: 3, AR033: 3, AR052: 3, AR243: 3, AR232: 3, AR248: 3, AR290: 3, AR089: 3, AR096: 3, AR238: 3, AR253: 3, AR234: 3, AR284: 3, AR229: 3, AR314: 2, AR295: 2, AR175: 2, AR316: 2, AR286: 2, AR289: 2, AR227: 2, AR247: 2, AR273: 2, AR296: 2, AR313: 2, AR182: 2, AR198: 2, AR231: 2, AR291: 2, AR060: 2,	1		
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18	HRABV49	988947	28	3 - 1757	204	<p>Thr-1 to Gly-7, Pro-35 to Gly-46, Ser-100 to Glu-111, Arg-144 to Pro-156, Gly-179 to His-187, Val-192 to Cys-201, Pro-254 to Asn-263, Lys-291 to Tyr-299, Gln-305 to Asn-312,</p>	<p>AR186: 2, AR237: 2, AR055: 2, AR226: 2, AR061: 2, AR184: 2, AR271: 2, AR177: 2, AR293: 2, AR292: 2, AR294: 2, AR219: 2, AR269: 2, AR246: 2, AR285: 2, AR218: 1, AR205: 1, AR233: 1, AR298: 1, AR274: 1, AR258: 1, AR256: 1, AR300: 1, AR259: 1, AR039: 1, AR240: 1, AR244: 1, AR185: 1, AR204: 1, AR241: 1, AR194: 0, AR179: 0, AR275: 0, AR206: 0, AR202: 0, AR192: 0, AR104: 0 H0597: 1, H0435: 1 and H0543: 1.</p>	11q12	<p>106100, 147050, 259700, 259770, 600045, 601884</p>
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Val-332 to Tyr-340, Pro-347 to Cys-357, Gly-384 to Trp-394, Pro-398 to Cys-403, Pro-406 to Trp-421, Cys-444 to Gln-455, Arg-457 to Ser-466, Pro-469 to Asp-480, Lys-488 to Asp-501, Ala-571 to His-576.	AR186: 1, AR166: 1, AR261: 1, AR312: 1, AR219: 1, AR296: 1, AR240: 1, AR313: 1, AR175: 1, AR222: 1, AR202: 1, AR289: 1, AR033: 1, AR253: 1, AR309: 1, AR177: 1, AR291: 1, AR234: 1, AR288: 1, AR266: 1, AR254: 1, AR267: 1, AR232: 1, AR252: 1, AR239: 1, AR089: 1, AR229: 1, AR218: 1, AR283: 1, AR242: 1, AR274: 1, AR262: 1, AR182: 1, AR295: 1, AR299: 1, AR200: 1, AR174: 1, AR236: 1, AR210: 1, AR183: 1, AR247: 1, AR277: 1, AR270: 1, AR292: 1, AR104: 1, AR096: 1, AR055: 1, AR179: 1, AR053: 1, AR316: 1, AR227: 1, AR251: 0, AR258: 0, AR275: 0, AR245: 0, AR265: 0, AR060: 0, AR294: 0, AR172: 0, AR314: 0,					
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AR241:	0, AR221:	0,						
AR233:	0, AR217:	0,						
AR162:	0, AR231:	0,						
AR185:	0, AR244:	0,						
AR268:	0, AR194:	0,						
AR061:	0, AR256:	0,						
AR171:	0, AR284:	0,						
AR271:	0, AR216:	0,						
AR211:	0, AR168:	0,						
AR205:	0, AR161:	0,						
AR198:	0, AR226:	0,						
AR039:	0, AR287:	0,						
AR196:	0, AR249:	0,						
AR293:	0, AR259:	0,						
AR281:	0, AR248:	0,						
AR308:	0, AR297:	0,						
AR207:	0, AR206:	0,						
AR164:	0, AR238:	0,						
AR243:	0, AR204:	0,						
AR176:	0, AR235:	0,						
AR315:	0, AR298:	0,						
AR311:	0, AR203:	0,						
AR300:	0, AR273:	0,						
AR169:	0, AR280:	0,						
AR285:	0, AR178:	0,						
AR191:	0, AR180:	0,						
AR201:	0, AR188:	0,						
AR237:	0, AR286:	0,						
AR189:	0, AR290:	0,						
AR255:	0, AR190:	0,						

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19	HNGGK47	993602	29	265 - 372	205	Trp-26 to Lys-36.	L0368: 1, L0789: 1, L0663: 1, H0519: 1, H0682: 1, H0670: 1, H0521: 1, H0522: 1, L0740: 1, L0779: 1, S0260: 1, H0667: 1 and H0543: 1. AR204: 21, AR205: 14, AR241: 10, AR243: 8, AR194: 8, AR206: 8, AR246: 7, AR244: 7, AR198: 7, AR186: 6, AR060: 6, AR271: 6, AR192: 6, AR104: 5, AR039: 5, AR240: 5, AR275: 5, AR313: 5, AR089: 5, AR273: 4, AR184: 4, AR033: 4, AR312: 4, AR055: 4, AR247: 4, AR177: 4, AR282: 4, AR299: 4, AR226: 4, AR053: 4, AR295: 4, AR061: 4, AR052: 4, AR182: 4, AR316: 4, AR293: 3, AR185: 3, AR202: 3, AR265: 3, AR274: 3, AR267: 3, AR096: 3, AR213: 3, AR296: 3, AR290: 3, AR183: 3,		
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20	HAJRC82	993759	30	2 - 2047	206	Asp-21 to Tyr-27, Pro-66 to Leu-72, Glu-99 to Ala-105,	AR309: 3, AR285: 3, AR229: 3, AR238: 3, AR294: 3, AR232: 3, AR277: 3, AR268: 2, AR269: 2, AR270: 2, AR249: 2, AR291: 2, AR300: 2, AR292: 2, AR263: 2, AR298: 2, AR289: 2, AR284: 2, AR259: 2, AR233: 2, AR175: 2, AR283: 2, AR231: 2, AR310: 2, AR227: 2, AR237: 2, AR234: 1, AR286: 1, AR219: 1, AR256: 1, AR266: 1, AR218: 1, AR258: 1, AR281: 1, AR179: 1, AR314: 1, AR280: 1, AR251: 1, AR248: 1, AR253: 1, AR315: 0 L0522: 3, L0748: 2, H0171: 1, H0686: 1, L0034: 1, H0615: 1, L0764: 1, L0794: 1, S0052: 1, S0374: 1 and L0601: 1.	9	

Gly-111 to Val-120, Gln-132 to Ile-138, Asp-152 to Ala-159, Lys-165 to Arg-170, Thr-222 to Cys-229, Arg-265 to Tyr-270, Ser-274 to Asp-283, Asp-299 to Ser-306, Val-316 to Arg-322, Asp-333 to Lys-346, Ser-447 to Arg-452, Asn-589 to Glu-594.	AR039: 2, AR052: 2, AR282: 2, AR213: 2, AR229: 2, AR234: 2, AR277: 2, AR186: 2, AR309: 1, AR267: 1, AR313: 1, AR291: 1, AR269: 1, AR312: 1, AR249: 1, AR253: 1, AR247: 1, AR270: 1, AR089: 1, AR061: 1, AR283: 1, AR284: 1, AR055: 1, AR175: 1, AR096: 1, AR060: 1, AR268: 1, AR316: 1, AR226: 1, AR240: 1, AR300: 1, AR033: 1, AR237: 1, AR298: 1, AR185: 1, AR244: 1, AR299: 1, AR281: 1, AR238: 1, AR273: 1, AR204: 1, AR246: 0, AR232: 0, AR182: 0, AR053: 0, AR177: 0, AR271: 0, AR256: 0, AR259: 0, AR295: 0, AR233: 0, AR194: 0, AR285: 0, AR274: 0, AR198: 0, AR266: 0, AR192: 0, AR251: 0, AR296: 0, AR218: 0.
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21	HJABW64	994001	31	1 - 465	207	Ala-26 to Thr-33, Ser-52 to Glu-58, Thr-83 to Leu-92.	L0366: 1, H0665: 1, S0242: 1, S0194: 1 and H0543: 1.	1		
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22	HAJBT01	997082	32	1290 - 2018	208	Gly-83 to Asp-93, Pro-106 to Val-113, Asp-123 to Val-141, Lys-143 to Ser-192, Ile-204 to Ser-209.	S0051: 1, H0038: 1, H0413: 1, T0069: 1, T0041: 1, H0494: 1, L0369: 1, L0794: 1, L0649: 1, L0803: 1, L0650: 1, L0651: 1, L0666: 1, H0520: 1, H0435: 1, H0658: 1, H0666: 1, H0214: 1, S0028: 1, L0439: 1, L0755: 1 and L0593: 1. AR217: 80, AR052: 80, AR280: 79, AR171: 75, AR314: 63, AR172: 61, AR169: 59, AR216: 57, AR168: 53, AR225: 53, AR270: 50, AR266: 50, AR223: 48, AR315: 47, AR218: 44, AR170: 44, AR214: 41, AR219: 39, AR224: 39, AR274: 39, AR291: 38, AR184: 36, AR290: 35, AR215: 34, AR222: 34, AR292: 34, AR268: 33, AR296: 33, AR210: 33, AR267: 32, AR269: 32, AR212: 31, AR183: 30, AR213: 30, AR173: 29, AR309: 28, AR312: 27, AR313: 27,		
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23	H7TSC95	997589	33	459 - I	209	Phe-28 to Tyr-35, Phe-59 to Trp-65, Gln-89 to Ser-98, Glu-127 to Glu-133.	H0444: 1, S0394: 1, L0596: 1, L0589: 1, L0594: 1, L0601: 1, S0011: 1, S0412: 1, S0462: 1 and S0460: 1, AR313: 8, AR274: 8, AR251: 8, AR266: 8, AR273: 7, AR218: 7, AR184: 7, AR219: 6, AR292: 6, AR096: 6, AR247: 6, AR271: 6, AR290: 6, AR243: 6, AR053: 6, AR175: 6, AR263: 6, AR183: 5, AR295: 5, AR280: 5, AR312: 5, AR315: 5, AR310: 5, AR240: 5, AR265: 4, AR293: 4, AR268: 4, AR309: 4, AR291: 4, AR177: 4, AR289: 4, AR256: 4, AR285: 4, AR286: 4, AR267: 4, AR284: 4, AR270: 4, AR269: 4, AR241: 4, AR182: 4, AR089: 4, AR052: 4, AR296: 4, AR316: 4, AR299: 4, AR314: 3, AR298: 3, AR282: 3, AR248: 3, AR213: 3,	7p14	10776, 138079, 138079, 139191, 180104, 203740, 601472, 601649
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24	HTEIS02	999204	34	2 - 373	210				AR310: 3, AR184: 3, AR246: 3, AR204: 2, AR205: 2, AR182: 2, AR274: 2, AR312: 2, AR295: 2, AR269: 1, AR247: 1, AR234: 1, AR194: 1, AR213: 1, AR061: 1, AR096: 1, AR293: 1, AR055: 1, AR277: 1, AR282: 1,	2q37.1	181031	

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AR244: 1, AR283: 1,					
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AR198: 0, AR177: 0,					
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AR175: 0, AR218: 0,					
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AR243: 0, AR231: 0,					
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25	HTXQD18	1000599	35	2 - 577	211	Asn-68 to Glu-73.	H0038: 3, H0587: 1, H0013: 1, L0118: 1 and H0024: 1. AR281: 9, AR205: 9, AR315: 8, AR283: 7, AR289: 7, AR204: 7, AR251: 7, AR055: 6, AR299: 6, AR247: 6, AR310: 6, AR033: 6, AR265: 6, AR266: 6, AR282: 6, AR202: 6, AR284: 6, AR060: 6, AR271: 6, AR246: 5, AR249: 5, AR243: 5, AR295: 5, AR280: 5, AR089: 5, AR241: 5, AR312: 5, AR263: 5, AR198: 5, AR277: 5, AR052: 5, AR206: 5, AR053: 5, AR194: 5, AR061: 5, AR316: 4, AR286: 4, AR213: 4, AR313: 4, AR291: 4, AR298: 4, AR039: 4, AR186: 4, AR273: 4, AR269: 4, AR183: 4, AR182: 4, AR270: 4, AR285: 4, AR309: 4, AR268: 4, AR232: 4, AR104: 4, AR294: 4,	9		
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77

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AR202: 1, AR207: 1,							
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27	HPMSA16	1011769	37	294 - 121	213		AR236: 0, AR174: 0, AR203: 0, AR289: 0, AR287: 0, AR280: 0, AR308: 0, AR195: 0, AR254: 0, AR237: 0, AR170: 0, AR222: 0, AR230: 0, AR235: 0, AR297: 0, AR163: 0, AR255: 0, AR193: 0 L0375: 1		
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28	HTTKE35	1027025	38	1 - 780	214	Gly-1 to His-9, Glu-50 to Pro-58, Ala-84 to Val-92, Ser-135 to Arg-152, Gly-185 to Ser-191, Arg-209 to Gly-215.	AR286: 0, AR213: 0, AR273: 0, AR249: 0, AR248: 0, AR299: 0, AR281: 0, AR313: 0, AR198: 0, AR182: 0, AR052: 0, AR300: 0, AR183: 0, AR179: 0, AR039: 0, AR231: 0, AR175: 0, AR234: 0, AR238: 0, AR259: 0, AR315: 0, AR293: 0, AR053: 0, AR240: 0, AR258: 0, AR284: 0, AR296: 0, AR265: 0, AR219: 0, AR104: 0, AR237: 0, AR268: 0, AR246: 0, AR280: 0, AR205: 0, AR291: 0, AR218: 0	15q26.1	166800, 210900
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AR285:	2, AR266:	2,
AR297:	2, AR234:	2,
AR277:	2, AR089:	2,
AR222:	2, AR200:	2,
AR096:	2, AR171:	2,
AR210:	2, AR292:	2,
AR286:	2, AR288:	2,
AR179:	2, AR232:	2,
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AR243:	1, AR215:	1,
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AR259:	1, AR260:	1,
AR258:	1, AR271:	1,
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AR219:	1, AR256:	1,
AR314:	1, AR280:	1,
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AR197:	1, AR218:	1,

29	HCRPE25	1034245	39	283 - 975	215	Lys-1 to Ser-6, Ser-81 to Asn-87.	AR241: 0, AR224: 0, AR170: 0, AR242: 0, AR281: 0, AR248: 0, AR245: 0, AR264: 0, AR194: 0, AR199: 0 H0265: 3, S0354: 1, H0634: 1, H0436: 1 and L0751: 1.	11p11-q11	133701, 171650, 176930, 176930, 600224, 600811
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30	HWAGX49	1034246	40	130 - 867	216	Cys-7 to His-15, Arg-17 to Thr-22, Ser-96 to Asn-102, Pro-170 to Gly-183.	AR268: 32, AR290: 24, AR270: 18, AR184: 14, AR267: 12, AR249: 9, AR269: 9, AR263: 8, AR182: 6, AR313: 4, AR175: 4, AR266: 4, AR096: 4, AR299: 4, AR234: 3, AR183: 3,		

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AR052:	2, AR202:	2,							
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AR298:	2, AR231:	2,							
AR277:	2, AR227:	2,							
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AR274:	0, AR294:	0,							

31	HTENW06	1040030	41	144 - 1145	217	Gly-5 to Glu-16, Gly-28 to Ser-34, Pro-186 to Pro-196, Ser-213 to Asn-219, Ser-249 to Ile-267.	AR233: 0, AR198: 0, AR275: 0, AR314: 0, AR219: 0, AR258: 0, AR204: 0, AR104: 0, AR206: 0 H0581: 2, L0755: 2, H0662: 1, S0358: 1, H0580: 1 and H0591: 1.	10q26.3	263700
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32	HODFW19	1052332	42	2 - 1249	218	Ala-22 to Thr-27, Asp-42 to Thr-49, Pro-77 to Ala-83,	AR282: 3, AR298: 3, AR263: 3, AR237: 3, AR310: 3, AR227: 3, AR243: 3, AR295: 3, AR240: 3, AR294: 3, AR290: 3, AR292: 3, AR268: 3, AR284: 2, AR291: 2, AR285: 2, AR249: 2, AR233: 2, AR232: 2, AR274: 2, AR283: 2, AR300: 2, AR253: 2, AR096: 2, AR231: 2, AR275: 2, AR258: 2, AR177: 2, AR234: 2, AR175: 2, AR259: 2, AR229: 1, AR256: 1, AR244: 1, AR219: 1, AR218: 1, AR179: 1, AR194: 1, AR315: 1, AR265: 0, AR281: 0, AR314: 0, AR280: 0 H0253: 7, L0439: 3, H0616: 2, H0617: 1, H0038: 1, H0560: 1, H0652: 1, L0774: 1, L0592: 1 and S0021: 1. AR238: 295, AR061: 198, AR237: 182, AR231: 170, AR226:		
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Ser-133 to Leu-140, Asp-152 to Ala-157.	144, AR227: 116, AR232: 104, AR234: 104, AR229: 102, AR233: 40, AR267: 35, AR089: 16, AR316: 14, AR271: 12, AR163: 7, AR202: 7, AR255: 5, AR186: 5, AR282: 5, AR263: 5, AR264: 5, AR055: 4, AR172: 4, AR206: 4, AR265: 4, AR194: 4, AR251: 3, AR308: 3, AR104: 3, AR171: 3, AR060: 3, AR192: 3, AR162: 3, AR249: 3, AR250: 3, AR246: 3, AR313: 3, AR274: 3, AR273: 2, AR052: 2, AR033: 2, AR257: 2, AR298: 2, AR185: 2, AR253: 2, AR184: 2, AR299: 2, AR270: 2, AR312: 2, AR240: 2, AR247: 2, AR207: 2, AR204: 2, AR096: 2, AR205: 2, AR179: 2, AR277: 2, AR295: 2, AR039: 2, AR164: 2, AR300: 2, AR243: 2, AR242: 2
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33	HACAD35	1052372	43	1 - 612	219	Lys-8 to Leu-16, Ala-47 to Gly-53, Gly-64 to Glu-72, Ala-93 to Gln-98.	AR184: 4, AR284: 4, AR292: 4, AR298: 4, AR263: 4, AR295: 4, AR182: 3, AR269: 3, AR175: 3, AR270: 3, AR293: 3, AR289: 3, AR285: 3, AR183: 3, AR267: 3, AR286: 3, AR296: 3, AR310: 2, AR194: 2, AR266: 2, AR268: 2, AR290: 2, AR291: 2, AR282: 2, AR198: 2, AR294: 2				

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AR277:	1, AR271:	1,
AR055:	1, AR052:	1,
AR061:	1, AR316:	1,
AR247:	1, AR259:	1,
AR226:	1, AR219:	1,
AR060:	1, AR177:	1,
AR246:	1, AR253:	1,
AR033:	1, AR234:	1,
AR089:	1, AR233:	1,
AR299:	1, AR258:	1,
AR186:	1, AR265:	1,
AR275:	1, AR096:	1,
AR256:	1, AR244:	1,
AR241:	1, AR300:	1,
AR227:	1, AR231:	1,
AR104:	1, AR232:	0,
AR249:	0, AR283:	0,
AR237:	0, AR218:	0,
AR243:	0, AR229:	0,
AR185:	0, AR309:	0,
AR281:	0, AR202:	0,
AR039:	0, AR273:	0,
AR240:	0, AR248:	0,
AR192:	0, AR315:	0,
AR251:	0, AR314:	0,
AR280:	0, AR206:	0,
AR204:	0, AR274:	0,
AR205:	0, AR213:	0,

34	HHPDX35	1056108	44	3 - 764	220	Gln-56 to Gln-61.	AR053: 0 L0637: 2, L0783: 2, L0777: 2, S6022: 1, H0392: 1, H0586: 1, H0050: 1, L0809: 1, L0759: 1 and S0192: 1. AR193: 442, AR228: 356, AR254: 313, AR288: 296, AR239: 294, AR210: 285, AR245: 265, AR287: 254, AR230: 251, AR199: 238, AR190: 228, AR203: 227, AR201: 227, AR242: 217, AR174: 211, AR211: 207, AR207: 198, AR260: 197, AR272: 196, AR163: 194, AR250: 193, AR262: 192, AR227: 191, AR255: 187, AR252: 184, AR236: 179, AR257: 179, AR180: 176, AR188: 174, AR195: 168, AR197: 168, AR200: 161, AR178: 161, AR191: 159, AR212: 156, AR189: 156,		
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35	HNFW14	1057738	45	2 - 400	221	Leu-9 to Met-15, Thr-25 to Cys-37, Glu-40 to Thr-46.	AR249: 0, AR052: 0 L0748: 4, L0105: 2, H0031: 2, H0494: 2, L0637: 2, L0662: 2, L0803: 2, L0666: 2, H0624: 1, H0171: 1, L0005: 1, L0717: 1, H0411: 1, H0587: 1, H0485: 1, H0105: 1, H0051: 1, H0622: 1, L0363: 1, L0364: 1, L0389: 1, L0659: 1, S0374: 1, H0670: 1, S0028: 1, L0743: 1, L0754: 1 and L0758: 1. AR248: 593, AR249: 232, AR253: 176, AR251: 161, AR219: 123, AR268: 122, AR315: 120, AR218: 118, AR314: 113, AR291: 105, AR183: 97, AR290: 91, AR270: 91, AR269: 81, AR247: 78, AR280: 76, AR281: 67, AR177: 65, AR298: 64, AR096: 64, AR295: 63, AR240: 60, AR182: 59, AR175: 54, AR234: 54, AR267: 54, AR179: 54,		
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36	HUKDS43	1061241	46	312 - 755	222	Pro-6 to Pro-29, Gly-67 to Gln-72, Cys-93 to Phe-100.	H0271: 2 AR273: 52, AR274: 49, AR243: 43, AR281: 37, AR053: 37, AR205: 36, AR052: 36, AR244: 35, AR312: 34, AR309: 34, AR271: 33, AR280: 33, AR206: 32, AR213: 32, AR246: 30, AR310: 30, AR315: 29, AR314: 23, AR263: 23, AR275: 22, AR096: 20, AR192: 20, AR039: 20, AR198: 19, AR218: 17, AR247: 17, AR313: 16, AR265: 16, AR219: 15, AR295: 15, AR316: 14, AR194: 14, AR299: 13, AR282: 13, AR089: 13, AR283: 12, AR202: 12, AR177: 12, AR175: 12, AR251: 11, AR291: 11, AR266: 11, AR240: 11, AR292: 11, AR241: 11, AR300: 10, AR256: 10, AR290: 9, AR284: 9, AR285: 9, AR185: 9, AR104: 9, AR183: 9, AR289: 8, AR060: 8, AR033: 8, AR298: 8, AR186: 8,		
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37	HAOAD02	1063218	47	354 - 3500	223	Ser-34 to His-40, Phe-104 to Glu-109, Asn-116 to Thr-127, Gly-138 to Glu-148, Gly-159 to Glu-165, Arg-192 to Ser-199, Ser-209 to Phe-215, Ser-242 to Ser-248.	H0135: 1, H0059: 1, S0344: 1, L0770: 1, L0766: 1, H0519: 1, S3014: 1, L0740: 1, L0752: 1, H0668: 1 and H0543: 1. AR222:1327, AR216:1210, AR210:1086, AR172:1078, AR224:1033, AR217: 987, AR221: 969, AR189: 810, AR190: 803, AR308: 729, AR218: 693, AR188: 645, AR211: 630, AR253: 609, AR191: 600, AR219: 592, AR212: 573, AR312: 554, AR173: 543, AR274: 537, AR168: 532, AR272: 530, AR179: 528, AR309: 506, AR171: 503, AR270: 493, AR250: 492, AR269: 489, AR280: 480, AR174: 478, AR170: 458, AR052: 454, AR176: 450, AR290: 442,	1q21	104770, 107670, 110700, 135940, 145001, 146790, 152445, 152445, 159001, 174000, 179755, 182860, 182860, 182860, 191315, 230800, 230800, 266200, 600897, 601105, 601412, 601652, 602491
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AR262: 263, AR310: 262, AR249: 259, AR162: 253, AR181: 251, AR165: 247, AR234: 245, AR184: 244, AR185: 244, AR282: 237, AR292: 235, AR265: 233, AR263: 230, AR300: 229, AR275: 226, AR230: 226, AR225: 224, AR192: 216, AR039: 215, AR237: 212, AR255: 209, AR196: 209, AR193: 202, AR231: 201, AR053: 200, AR277: 188, AR266: 183, AR281: 172, AR251: 169, AR226: 165, AR289: 164, AR291: 164, AR296: 163, AR295: 160, AR259: 159, AR287: 153, AR238: 150, AR186: 149, AR197: 145, AR294: 143, AR243: 135, AR201: 129, AR232: 129, AR229: 128, AR242: 125,								
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38	HOUCD83	1070724	48	3892 - 2978	224			AR285: 125, AR298: 123, AR239: 123, AR195: 122, AR241: 115, AR246: 115, AR033: 110, AR198: 105, AR258: 105, AR257: 93, AR244: 93, AR260: 92, AR261: 92, AR104: 84, AR061: 73, AR204: 72, AR206: 69, AR235: 67, AR194: 66, AR286: 61, AR227: 59, AR233: 58, AR283: 56, AR202: 55, AR228: 40, AR055: 37, AR207: 32 L0759: 3, S0314: 2, L0744: 2, L0756: 2, L0755: 2, H0391: 1, H0052: 1, H0050: 1, S0318: 1, S0338: 1 and H0144: 1.			AR089: 391, AR060: 321, AR104: 230, AR316: 189, AR185: 142, AR299: 68, AR283: 65, AR096: 58, AR282: 46, AR240: 42, AR277: 38, AR313: 37, AR230: 23, AR228: 22, AR287: 21, AR239: 19, AR297:	3q13	146200, 190300, 258900, 600882
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9, AR189: 9, AR246:					
9, AR175: 9, AR199:					
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8, AR295: 8, AR226:					
8, AR256: 8, AR214:					
8, AR172: 8, AR052:					
8, AR216: 8, AR198:					
8, AR201: 8, AR221:					
8, AR285: 7, AR269:					
7, AR195: 7, AR206:					
7, AR171: 7, AR310:					
7, AR258: 7, AR273:					
7, AR168: 7, AR275:					
7, AR309: 7, AR210:					
7, AR274: 7, AR244:					
7, AR270: 7, AR053:					
7, AR183: 6, AR250:					
6, AR245: 6, AR213:					
6, AR251: 6, AR193:					
6, AR247: 6, AR192:					
6, AR241: 6, AR222:					
6, AR268: 6, AR290:					
6, AR218: 6, AR311:					
5, AR039: 5, AR205:					
5, AR219: 5, AR271:					
5, AR264: 5, AR312:					
5, AR170: 4, AR253:					
4, AR308: 4, AR196:					
4, AR211: 3, AR259:					

39	HOFMS43	1071027	49	25 - 594	225	<p>Arg-9 to Asp-30, Pro-74 to Asn-79, Tyr-97 to Tyr-103, Ser-116 to Pro-129, Pro-144 to Glu-155.</p>	<p>3, AR194: 3, AR248: 3, AR292: 3, AR265: 3, AR263: 3, AR249: 2, AR314: 2, AR169: 2, AR280: 2, AR298: 2, AR284: 1, AR281: 1, AR315: 1, AR184: 1 L0754: 4, S0040: 1, L0809: 1, H0672: 1, L0777: 1 and L0780: 1.</p>		
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AR213:	7, AR229:	7,
AR238:	7, AR310:	7,
AR231:	6, AR089:	6,
AR246:	6, AR267:	6,
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AR192:	6, AR299:	6,
AR263:	6, AR052:	6,
AR175:	6, AR039:	6,
AR234:	6, AR273:	5,
AR253:	5, AR274:	5,
AR277:	5, AR219:	5,
AR180:	5, AR300:	5,
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AR271:	5, AR204:	5,
AR237:	5, AR186:	4,
AR218:	4, AR240:	4,
AR053:	4, AR177:	4,
AR168:	4, AR060:	4,
AR312:	4, AR309:	4,
AR198:	4, AR185:	4,
AR232:	4, AR275:	4,
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AR178:	2, AR224:	2,
AR163:	2, AR254:	2,
AR162:	2, AR161:	2,
AR171:	2, AR242:	2,
AR264:	2, AR223:	2,

40	HMTAK58	1071812	50	2 - 553	226	Asn-55 to Pro-60, Ser-83 to Asp-100, Tyr-118 to His-124, Ser-129 to Trp-136.	AR222: 2, AR165: 2, AR225: 2, AR288: 2, AR166: 2, AR181: 1, AR212: 1, AR164: 1, AR287: 1, AR195: 1, AR172: 1, AR203: 1, AR211: 1, AR188: 1, AR196: 1, AR176: 1, AR245: 1, AR189: 1, AR190: 1, AR236: 1, AR215: 1, AR191: 1, AR169: 1, AR255: 1, AR252: 1, AR235: 1, AR214: 1, AR257: 1, AR207: 1, AR239: 0, AR200: 0, AR216: 0, AR170: 0, AR297: 0, AR193: 0, AR272: 0, AR250: 0, AR260: 0, AR261: 0, AR230: 0, AR221: 0, AR210: 0, AR174: 0, AR201: 0, AR197: 0, AR228: 0, AR199: 0, AR308: 0, AR311: 0, AR262: 0 H0415: 1	12p13	103950, 120580, 131440, 139130,
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41	H7TBB38	1073467	51	3 - 713	227			AR061: 1, AR248: 1, AR274: 1, AR204: 0, AR283: 0, AR202: 0, AR194: 0, AR192: 0, AR241: 0, AR198: 0, AR206: 0, AR055: 0, AR039: 0, AR273: 0, AR246: 0, AR275: 0, AR104: 0 H0518: 3, H0656: 2, H0341: 2, H0038: 2, H0616: 2, L0766: 2, L0748: 2, H0542: 2, H0543: 2, H0221: 1, H0158: 1, H0036: 1, H0571: 1, H0109: 1, H0634: 1, H0560: 1, H0561: 1, L0763: 1, L0662: 1, L0794: 1, L0775: 1, L0655: 1, L0663: 1, H0648: 1, H0525: 1 and H0423: 1.		
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AR268:	1, AR061:	1,						
AR198:	1, AR313:	1,						
AR096:	1, AR293:	1,						
AR089:	1, AR296:	1,						
AR248:	1, AR213:	1,						
AR227:	1, AR277:	1,						
AR238:	1, AR263:	1,						
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AR182:	1, AR300:	1,						
AR285:	1, AR270:	1,						
AR237:	1, AR251:	1,						
AR055:	1, AR244:	1,						
AR292:	1, AR271:	1,						
AR192:	1, AR233:	1,						
AR104:	1, AR184:	1,						
AR033:	1, AR234:	1,						
AR256:	1, AR259:	1,						
AR299:	1, AR265:	1,						
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AR289:	0, AR253:	0,						
AR274:	0, AR258:	0,						
AR283:	0, AR226:	0,						
AR206:	0, AR315:	0,						
AR241:	0, AR205:	0,						
AR202:	0, AR243:	0,						

42	HPWBS63	1077575	52	3 - 833	228	Arg-56 to Leu-61, Asp-77 to His-91, Lys-110 to Lys-115, Val-158 to Gln-177, Asp-186 to Thr-193, Lys-257 to Leu-269.	AR280: 0, AR219: 0, AR177: 0, AR232: 0, AR179: 0, AR039: 0, AR052: 0, AR053: 0, AR204: 0, AR281: 0, AR218: 0 S0198: 55, S0198: 55, S0274: 12, S0274: 12, S0252: 4, S0252: 4, S0270: 3, S0270: 3, S0264: 1, S0264: 1, S0268: 1, S0268: 1, S0228: 1 and S0228: 1. AR055: 17, AR104: 16, AR251: 15, AR186: 14, AR182: 13, AR061: 12, AR060: 12, AR286: 11, AR267: 10, AR258: 10, AR033: 10, AR310: 10, AR184: 10, AR298: 10, AR185: 9, AR285: 9, AR089: 9, AR284: 9, AR309: 8, AR270: 8, AR265: 8, AR280: 8, AR204: 8, AR052: 8, AR266: 8, AR053: 8, AR299: 8, AR316: 7, AR271: 7, AR253: 7, AR183: 7, AR289: 7, AR256: 7, AR259: 7,		

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43	HSDJH63	1078285	53	153 - 1358	229	Thr-17 to Lys-23, Lys-93 to Arg-98.	L0770: 3, L0794: 3, L0754: 3, H0266: 2, L0761: 2, L0774: 2, L0806: 2, L0439: 2, S0040: 1, H0484: 1, H0254: 1, H0255: 1, H0600: 1, S0210: 1, L0800: 1, L0764: 1, L0771: 1, L0773: 1, L0662: 1, L0775: 1, L0652: 1, L0665: 1, S0044: 1, L0745: 1, L0750: 1, L0786: 1 and L0757: 1. AR194: 31, AR206: 30, AR202: 30, AR039: 28, AR205: 27, AR246: 26, AR310: 25, AR265: 24, AR263: 23, AR289: 23, AR244: 22, AR266: 20, AR053: 20, AR284: 20, AR309: 19, AR241: 19, AR271: 19, AR282: 18, AR213: 18, AR243: 18, AR204: 18, AR312: 17, AR198: 17, AR052: 16, AR283: 16, AR251: 16, AR281: 16, AR192: 16, AR275: 16, AR277: 16, AR273: 16, AR315: 16,		
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115

44	HPLBQ44	1081779	54	3 - 986	230	HO657: 1, LO791: 1, HO689: 1, SO152: 1, SO260: 1 and HO445: 1. AR201: 1719, AR207: 999, AR197: 799, AR195: 698, AR204: 510, AR271: 496, AR192: 453, AR193: 431, AR212: 415, AR198: 377, AR163: 357, AR245: 319, AR205: 312, AR272: 310, AR166: 309, AR242: 297, AR254: 295, AR252: 294, AR277: 273, AR055: 272, AR308: 259, AR165: 256, AR161: 255, AR104: 255, AR250: 255, AR275: 251, AR185: 248, AR311: 239, AR162: 234, AR264: 208, AR060: 185, AR164: 177, AR274: 175, AR246: 173, AR053: 166, AR309: 159, AR039: 144, AR253: 132, AR033: 119, AR213: 116, AR312:	10		
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45	HAHBE26	1082286	55	43 - 771	231	His-8 to Gly-18, Tyr-109 to Lys-115, Asp-171 to Tyr-176.	S0036: 1, H0551: 1, H0264: 1, H0059: 1, H0494: 1, H0633: 1, S0144: 1, H0529: 1, H0666: 1, H0555: 1, S0390: 1, L0758: 1, S0031: 1 and H0543: 1. AR289: 34, AR281: 24, AR296: 23, AR265: 23, AR284: 22, AR285: 22, AR291: 22, AR266: 22, AR298: 21, AR315: 21, AR263: 20, AR286: 19, AR310: 19, AR292: 18, AR295: 17, AR258: 17, AR293: 17, AR283: 17, AR314: 16, AR280: 16, AR294: 16, AR309: 16, AR055: 14, AR312: 14, AR033: 14, AR268: 13, AR247: 13, AR282: 13, AR256: 13, AR194: 13, AR089: 12, AR269: 12, AR039: 12, AR202: 12, AR277: 12, AR205: 12, AR270: 12, AR299: 12, AR183: 12, AR316: 12, AR206: 12, AR182: 11, AR053: 11, AR259: 11, AR251: 11, AR313: 11,		
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46	HMMCH04	1082413	56	594 - 490	232	Gly-24 to Ala-30.	AR232: 11, AR096: 10, AR267: 10, AR244: 10, AR290: 10, AR300: 10, AR177: 10, AR246: 10, AR052: 10, AR213: 10, AR238: 9, AR175: 9, AR253: 9, AR229: 9, AR184: 9, AR243: 9, AR198: 8, AR227: 8, AR061: 8, AR226: 8, AR271: 8, AR204: 8, AR234: 8, AR192: 8, AR248: 8, AR233: 7, AR060: 7, AR241: 7, AR231: 7, AR218: 7, AR275: 6, AR249: 6, AR273: 6, AR219: 6, AR237: 6, AR186: 6, AR185: 6, AR240: 5, AR274: 4, AR179: 3, AR104: 3 H0002: 1 and S0364: 1. AR241: 172, AR198: 94, AR192: 89, AR313: 85, AR039: 73, AR219: 68, AR204: 64, AR271: 64, AR240: 60, AR229: 57, AR194: 54, AR218: 54, AR258: 53, AR089: 49, AR274: 49, AR096: 48,		
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47	HOFMN17	1083062	57	3 - 746	233	Pro-13 to Ile-21, Lys-77 to Arg-88.	AR298: 16, AR290: 16, AR266: 15, AR309: 15, AR248: 14, AR284: 14, AR283: 14, AR289: 14, AR253: 12, AR055: 8, AR061: 6 L0717: 1, H0581: 1, L0752: 1 and H0444: 1. AR312: 55, AR309: 45, AR281: 44, AR265: 39, AR310: 39, AR315: 38, AR280: 37, AR274: 35, AR053: 34, AR263: 34, AR314: 32, AR096: 31, AR194: 30, AR313: 30, AR213: 30, AR251: 29, AR052: 29, AR299: 27, AR089: 25, AR039: 23, AR273: 23, AR247: 22, AR202: 22, AR243: 21, AR205: 21, AR295: 21, AR282: 21, AR253: 21, AR283: 20, AR266: 20, AR246: 20, AR271: 20, AR206: 20, AR316: 20, AR244: 19, AR033: 18, AR300: 18, AR275: 17, AR192: 17, AR277: 16, AR292: 16, AR198: 16, AR270: 16, AR240: 16.				
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48	HKABY22	1083735	58	28 - 723	234	<p> AR241: 15, AR104: 15, AR289: 14, AR284: 14, AR285: 13, AR290: 13, AR232: 13, AR293: 13, AR185: 13, AR291: 13, AR204: 12, AR248: 12, AR177: 12, AR060: 12, AR218: 12, AR183: 11, AR268: 11, AR219: 11, AR186: 11, AR267: 11, AR294: 11, AR249: 11, AR184: 11, AR182: 10, AR296: 10, AR269: 10, AR231: 10, AR055: 10, AR286: 9, AR175: 9, AR238: 9, AR229: 8, AR298: 8, AR227: 8, AR061: 7, AR179: 7, AR237: 7, AR256: 6, AR226: 6, AR234: 6, AR233: 6, AR258: 4, AR259: 4 H0519: 3, H0624: 1, H0415: 1, H0592: 1, H0618: 1, H0561: 1 and L0366: 1. </p>	<p> AR241: 15, AR104: 15, AR289: 14, AR284: 14, AR285: 13, AR290: 13, AR232: 13, AR293: 13, AR185: 13, AR291: 13, AR204: 12, AR248: 12, AR177: 12, AR060: 12, AR218: 12, AR183: 11, AR268: 11, AR219: 11, AR186: 11, AR267: 11, AR294: 11, AR249: 11, AR184: 11, AR182: 10, AR296: 10, AR269: 10, AR231: 10, AR055: 10, AR286: 9, AR175: 9, AR238: 9, AR229: 8, AR298: 8, AR227: 8, AR061: 7, AR179: 7, AR237: 7, AR256: 6, AR226: 6, AR234: 6, AR233: 6, AR258: 4, AR259: 4 H0519: 3, H0624: 1, H0415: 1, H0592: 1, H0618: 1, H0561: 1 and L0366: 1. </p>		
						<p> His-1 to Gly-12, Pro-41 to Arg-47, Gln-52 to Thr-57, Leu-97 to Ala-103, </p>	<p> AR194: 3, AR184: 3, AR282: 2, AR310: 2, AR241: 1, AR269: 1, AR295: 1, AR265: 1, </p>		

Asp-131 to Asn-137, Gln-156 to Gly-161, Arg-170 to Arg-175, Pro-224 to His-232.	AR313: 1, AR204: 1, AR175: 1, AR284: 1, AR182: 1, AR267: 1, AR096: 1, AR053: 1, AR268: 1, AR266: 1, AR277: 1, AR052: 1, AR294: 1, AR183: 1, AR312: 1, AR039: 1, AR251: 1, AR232: 1, AR296: 1, AR089: 1, AR281: 1, AR238: 1, AR298: 1, AR286: 1, AR234: 1, AR247: 1, AR186: 1, AR104: 1, AR061: 1, AR270: 1, AR316: 0, AR285: 0, AR244: 0, AR314: 0, AR291: 0, AR246: 0, AR185: 0, AR033: 0, AR248: 0, AR280: 0, AR271: 0, AR192: 0, AR229: 0, AR263: 0, AR219: 0, AR055: 0, AR249: 0, AR315: 0, AR243: 0, AR299: 0, AR237: 0, AR253: 0, AR256: 0, AR259: 0, AR218: 0, AR202: 0, AR283: 0, AR177: 0, AR275: 0, AR289: 0.
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49	HLDOJ74	1083901	59	11 - 1939	235	Arg-9 to Val-17, Asn-87 to Val-100, Arg-175 to Ile-181, Gln-249 to Pro-258, Asp-271 to Asn-276, Arg-297 to Cys-303, Val-379 to Asp-393.	AR293: 0, AR300: 0, AR213: 0, AR233: 0, AR309: 0, AR240: 0, AR274: 0, AR206: 0, AR227: 0, AR060: 0, AR198: 0, AR179: 0, AR290: 0, AR205: 0, AR292: 0, AR258: 0, AR231: 0 L0743: 6, H0545: 2, H0676: 1, H0587: 1, H0494: 1, L0640: 1, L0769: 1, L0650: 1 and L0758: 1.		
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50	HTLDS55	1083991	60	1 - 1341	236		AR186: 6, AR246: 5, AR052: 4, AR251: 4, AR282: 4, AR055: 4, AR225: 4, AR202: 4, AR061: 4, AR309: 4, AR289: 3, AR182: 3, AR206: 3, AR060: 3, AR273: 3, AR291: 3, AR311: 3, AR310: 3, AR181: 3, AR265: 3, AR272: 3, AR284: 3, AR253: 3, AR053: 3, AR170: 3, AR298: 3, AR283: 3, AR162: 3, AR161: 3, AR296: 3, AR249: 3, AR171: 3, AR163: 3, AR089: 3, AR165: 3, AR192: 2, AR164: 2, AR241: 2, AR270: 2, AR221: 2, AR166: 2, AR223: 2, AR266: 2, AR292: 2, AR033: 2, AR269: 2, AR183: 2, AR312: 2,			

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AR299:	2, AR290:	2,							
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AR184:	2, AR263:	2,							
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AR237:	2, AR200:	2,							
AR195:	2, AR295:	2,							
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AR226:	1, AR177:	1,							
AR236:	1, AR174:	1,							
AR201:	1, AR231:	1,							
AR243:	1, AR287:	1,							
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AR232:	1, AR259:	1,							
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AR238:	1, AR258:	1,							

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51	HDQ GK06	1084465	61	225 - 2012	237	Arg-9 to Arg-16, Ala-55 to Lys-70, Trp-80 to Leu-93, Ser-120 to Asp-141,	AR184: 7, AR162: 6, AR161: 6, AR163: 6, AR310: 6, AR165: 5, AR166: 5, AR164: 5,					

Thr-148 to Glu-153, Arg-193 to Trp-216, Thr-221 to Ser-238, Gln-254 to Lys-264, Ser-302 to Ser-307, Gly-382 to Asp-389, Asn-415 to Ser-428, Tyr-444 to Asn-451, Asp-461 to Asn-474, Cys-504 to Asp-512, Cys-517 to His-523, Pro-525 to Gly-533, Pro-535 to Gln-540, Tyr-551 to Trp-559.	AR313: 5, AR308: 5, AR169: 5, AR263: 5, AR089: 4, AR172: 4, AR183: 4, AR229: 4, AR212: 4, AR265: 4, AR270: 4, AR196: 4, AR173: 4, AR299: 4, AR178: 4, AR238: 4, AR242: 4, AR171: 4, AR311: 4, AR312: 4, AR182: 4, AR231: 4, AR252: 4, AR215: 3, AR225: 3, AR096: 3, AR226: 3, AR269: 3, AR180: 3, AR280: 3, AR300: 3, AR297: 3, AR309: 3, AR216: 3, AR053: 3, AR213: 3, AR290: 3, AR272: 3, AR234: 3, AR314: 3, AR174: 3, AR236: 3, AR291: 3, AR247: 3, AR060: 3, AR296: 3, AR267: 3, AR315: 3, AR241: 3, AR240: 3, AR181: 3, AR175: 3, AR237: 3, AR266: 3, AR316: 3, AR264: 2, AR293: 2, AR251: 2, AR282: 2, AR298: 2,
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52	HNGME19	1084997	62	3 - 1187	238	Asp-5 to Gln-13, Gln-40 to Lys-48, Asp-57 to Ser-65, Asp-75 to Asp-87, Asp-111 to Gly-116, Leu-144 to Ser-151, Arg-231 to Glu-237, Ile-241 to Gly-246, Asp-270 to Leu-277, Thr-290 to Ser-295, Leu-301 to Glu-306.	L0657: 1, L0659: 1, L0809: 1, L0789: 1, L0666: 1, L0664: 1, H0144: 1, S0328: 1, L0754: 1, L0747: 1, L0731: 1, L0759: 1, H0445: 1, L0592: 1, S0026: 1, H0423: 1 and H0352: 1.	13q21	277900, 602085
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53	HELGO48	1085552	63	1 - 1296	239	Phe-6 to Pro-13.	AR237: 3, AR300: 3, AR251: 3, AR249: 3, AR299: 3, AR089: 3, AR296: 3, AR233: 3, AR227: 3, AR290: 3, AR270: 3, AR313: 3, AR309: 3, AR292: 3, AR295: 3, AR232: 3, AR229: 3, AR268: 3, AR284: 3, AR267: 2, AR277: 2, AR231: 2, AR285: 2, AR274: 2, AR298: 2, AR184: 2, AR259: 2, AR266: 2, AR256: 2, AR258: 1, AR175: 1, AR294: 1, AR286: 1, AR219: 1, AR218: 1, AR177: 1, AR265: 1, AR281: 1, AR291: 1, AR183: 1, AR179: 1, AR314: 1, AR234: 0, AR280: 0, AR315: 0 S0010: 4, L0471: 2, L0748: 2, S0031: 2, H0369: 1, H0263: 1, N0006: 1, H0373: 1, H0356: 1, H0644: 1, S0428: 1 and L0759: 1.	AR269: 13, AR241: 12, Xq21.33-	300046,
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AR274: 8, AR055: 8,		301835,
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AR238: 4, AR183: 4,		

54	HRDAF83	1085723	64	992 - 696	240	Asn-66 to Val-71, Glu-82 to Arg-90.	AR277: 4, AR175: 4, AR177: 4, AR299: 4, AR296: 4, AR313: 4, AR237: 4, AR291: 4, AR253: 4, AR247: 4, AR096: 3, AR226: 3, AR104: 3, AR232: 3, AR219: 3, AR218: 3, AR249: 2, AR259: 1, AR258: 1, AR256: 1, AR280: 0, AR315: 0, AR248: 0, AR314: 0 L0770: 4, T0049: 2, H0544: 2, H0268: 2, L0759: 2, H0662: 1, S0376: 1, S0045: 1, S0132: 1, H0587: 1, H0574: 1, H0599: 1, H0052: 1, H0546: 1, H0124: 1, H0163: 1, H0551: 1, H0269: 1, H0646: 1, L0631: 1, L0662: 1, L0653: 1, L0659: 1, L0666: 1, L0665: 1, S0037: 1, L0758: 1, H0595: 1, L0601: 1 and H0506: 1. AR241: 13, AR313: 9, AR244: 9, AR194: 8, AR186: 7, AR229: 7,	15q22.3- q23	118485, 151670, 231680,
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AR162: 6, AR164: 6,	601780
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AR192: 6, AR196: 6,	
AR163: 6, AR053: 6,	
AR202: 6, AR271: 6,	
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AR247: 6, AR178: 6,	
AR039: 6, AR294: 5,	
AR269: 5, AR207: 5,	
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AR246: 5, AR261: 5,	
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AR300: 5, AR206: 5,	
AR257: 5, AR262: 5,	
AR089: 5, AR060: 5,	
AR251: 5, AR181: 5,	
AR248: 5, AR185: 5,	
AR226: 5, AR274: 5,	
AR264: 5, AR237: 5,	
AR055: 5, AR275: 5,	
AR270: 5, AR033: 5,	
AR243: 5, AR296: 5,	
AR286: 5, AR292: 5,	
AR061: 4, AR182: 4,	
AR191: 4, AR096: 4,	

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55	HBJHS02	1085743	65	1 - 567	241	Ala-39 to Gly-44, Ala-89 to Tyr-97, Thr-103 to Gln-114, Pro-133 to Ser-138, Thr-143 to Pro-149.			AR213: 16, AR052: 13, 1 AR053: 12, AR194: 5, AR096: 5, AR061: 4, AR244: 4, AR263: 4, AR186: 3, AR204: 3, AR060: 3, AR251: 3, AR226: 3, AR241: 3,				

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56	HE8PB92	1086828	66	2 - 697	242	Lys-55 to Val-67.	AR226: 15, AR232: 7, AR227: 7, AR237: 6, AR238: 6, AR284: 5, AR269: 5, AR183: 4, AR293: 4, AR270: 4, AR266: 4, AR295: 4, AR292: 4, AR291: 3, AR219: 3, AR294: 3, AR258: 3, AR286: 3, AR218: 3, AR265: 3,			

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AR060: 1, AR248: 1,									
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AR243: 0, AR273: 0,									

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57	HTEMU66	1086856	67	462 - 962	243	His-3 to His-10, Thr-20 to Trp-26.	L0375: 1, L0666: 1, L0665: 1, L0352: 1, S0126: 1, H0648: 1, H0672: 1, L0750: 1, L0756: 1, L0779: 1, L0777: 1, L0752: 1 and L0755: 1. AR202: 11, AR206: 11, AR194: 11, AR284: 10, AR244: 9, AR289: 9, AR296: 9, AR205: 9, AR246: 9, AR052: 9, AR263: 9, AR198: 8, AR285: 8, AR266: 8, AR291: 8, AR295: 8, AR204: 7, AR192: 7, AR241: 7, AR053: 7, AR182: 7, AR039: 7, AR275: 7, AR310: 6, AR265: 6, AR269: 6, AR243: 6, AR292: 6, AR282: 6, AR298: 6, AR213: 6, AR309: 6, AR268: 6, AR251: 6, AR055: 6, AR232: 6, AR290: 6, AR312: 6, AR271: 5, AR060: 5, AR270: 5, AR033: 5, AR227: 5, AR316: 5, AR277: 5, AR293: 5,		
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58	HE2PQ73	1089285	68	127 - 1020	244	Thr-28 to Ser-33.	AR286: 5, AR231: 5, AR294: 5, AR273: 5, AR313: 5, AR184: 5, AR253: 5, AR061: 5, AR226: 5, AR267: 5, AR186: 5, AR177: 5, AR089: 5, AR185: 5, AR281: 5, AR237: 4, AR234: 4, AR300: 4, AR274: 4, AR238: 4, AR299: 4, AR249: 4, AR104: 4, AR233: 4, AR315: 4, AR247: 4, AR175: 4, AR229: 4, AR280: 4, AR240: 4, AR248: 3, AR096: 3, AR314: 3, AR283: 3, AR218: 3, AR183: 3, AR219: 3, AR258: 3, AR256: 2, AR179: 2, AR259: 2 H0616: 1	14q24-q32	107970, 115650, 123270, 182600, 245200, 251600, 270100, 276900,
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59	HOHEC71	1090538	69	42 - 545	245	Val-8 to Ser-18, Pro-35 to Gln-43, Gly-60 to Tyr-66, Ala-74 to Lys-80, Ser-90 to Arg-97, Ser-155 to Ala-163.	AR277: 3, AR061: 3, AR234: 3, AR202: 3, AR237: 3, AR231: 3, AR204: 3, AR227: 2, AR233: 2, AR281: 2, AR246: 1 L0439: 3, L0748: 2, H0624: 1, H0171: 1 and H0623: 1.		
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60	HCE3J64	1090546	70	3 - 692	246	Thr-1 to Lys-8, Lys-26 to Asp-33, Gln-60 to Trp-70, Phe-99 to Lys-106, Asp-126 to Asn-137,	H0485: 1, H0427: 1, S0280: 1, L0021: 1, T0082: 1, H0618: 1, S0010: 1, H0007: 1, H0085: 1, H0235: 1, H0327: 1, H0545: 1, H0081: 1, H0012: 1, H0082: 1, H0020: 1, H0051: 1, S6028: 1, H0286: 1, H0424: 1, H0418: 1, H0628: 1, H0124: 1, S0036: 1, H0040: 1, H0063: 1, H0116: 1, H0132: 1, H0652: 1, L0763: 1, L0646: 1, L0794: 1, L0766: 1, L0650: 1, L0651: 1, L0653: 1, L0792: 1, L0666: 1, L0663: 1, L0665: 1, H0144: 1, S0380: 1, L0754: 1, L0755: 1, S0031: 1, L0608: 1, L0366: 1, H0653: 1 and H0352: 1.	3q26.1- q26.33	138160, 138160, 177400, 600044
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Arg-139 to Glu-145, Met-157 to Tyr-164, Asn-167 to Arg-174, Lys-200 to Leu-207.	AR192: 53, AR241: 52, AR265: 45, AR271: 44, AR310: 38, AR263: 37, AR213: 37, AR274: 36, AR282: 35, AR052: 34, AR273: 34, AR309: 33, AR300: 31, AR053: 31, AR275: 31, AR281: 30, AR295: 30, AR232: 29, AR289: 28, AR104: 28, AR299: 27, AR096: 27, AR266: 27, AR312: 27, AR316: 27, AR033: 26, AR240: 26, AR277: 25, AR283: 24, AR284: 23, AR247: 23, AR285: 23, AR251: 22, AR313: 22, AR060: 22, AR089: 22, AR177: 21, AR291: 21, AR315: 21, AR292: 20, AR286: 20, AR280: 19, AR186: 19, AR185: 19, AR270: 19, AR298: 18, AR296: 18, AR055: 18, AR227: 18, AR269: 18, AR238: 17, AR182: 17, AR175: 17, AR294: 17, AR268: 17, AR231: 16, AR314: 16, AR226: 16, AR218: 16, AR234: 16,					
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61	HCE3L81	1090711	71	1 - 675	247	Pro-1 to Ala-9, Ser-17 to Ser-24, Pro-48 to Gln-53, Asn-100 to Ala-107, Ser-127 to Ser-132.	AR253: 16, AR293: 14, AR256: 14, AR229: 14, AR183: 14, AR061: 14, AR219: 14, AR290: 14, AR267: 13, AR258: 13, AR249: 13, AR237: 13, AR233: 13, AR184: 12, AR259: 11, AR248: 10, AR179: 9 H0052: 2, L0759: 2, S0376: 1, H0333: 1, S0388: 1, L0637: 1, L0636: 1, L0742: 1 and L0439: 1.	AR274: 27, AR251: 21, AR052: 21, AR309: 20, AR273: 20, AR243: 19, AR313: 19, AR053: 19, AR312: 18, AR271: 18, AR266: 17, AR310: 17, AR213: 15, AR186: 15, AR265: 14, AR198: 13, AR275: 12, AR060: 12, AR089: 11, AR291: 11, AR290: 11, AR241: 11, AR185: 11, AR289: 11, AR104: 11, AR270: 11, AR285: 11, AR253: 10, AR240: 10, AR192: 10, AR269: 10, AR316: 9,	17		
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62	HUFDJ62	1090753	72	26 - 667	248	Leu-3 to Ala-20.	AR282: 9, AR268: 9, AR286: 9, AR247: 9, AR096: 9, AR033: 9, AR182: 9, AR205: 9, AR263: 9, AR184: 9, AR175: 9, AR295: 8, AR299: 8, AR296: 8, AR204: 8, AR293: 8, AR267: 8, AR194: 8, AR284: 8, AR248: 8, AR238: 8, AR055: 8, AR292: 8, AR061: 7, AR283: 7, AR298: 7, AR039: 7, AR206: 7, AR300: 7, AR202: 6, AR256: 6, AR277: 6, AR183: 6, AR226: 6, AR237: 6, AR232: 6, AR229: 6, AR218: 6, AR231: 6, AR246: 6, AR177: 5, AR294: 5, AR219: 5, AR244: 5, AR234: 5, AR249: 5, AR233: 4, AR227: 4, AR258: 4, AR259: 4, AR179: 3, AR281: 1, AR280: 1, AR315: 0 H0052: 2, H0663: 1 and H0555: 1.		
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				Gly-89 to Pro-98, Val-111 to Ile-124, Glu-138 to Tyr-143.	AR194: 24, AR202: 22, AR263: 22, AR280: 20, AR246: 20, AR314: 19, AR244: 18, AR283: 18, AR265: 16, AR310: 15, AR206: 14, AR033: 13, AR241: 13, AR096: 13, AR282: 13, AR205: 12, AR295: 12, AR192: 12, AR243: 12, AR053: 12, AR198: 12, AR299: 11, AR300: 11, AR277: 11, AR266: 11, AR247: 11, AR213: 10, AR312: 10, AR284: 10, AR183: 10, AR089: 10, AR309: 10, AR251: 10, AR316: 10, AR039: 10, AR055: 9, AR052: 9, AR285: 9, AR313: 9, AR292: 9, AR177: 9, AR289: 8, AR271: 8, AR232: 8, AR294: 7, AR229: 7, AR248: 7, AR175: 7, AR227: 7, AR298: 7, AR184: 7, AR253: 7, AR204: 7, AR286: 7, AR270: 7, AR234: 7, AR226: 6, AR268: 6, AR231: 6, AR238: 6.		
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63	HE8FC96	1090865	73	75 - 989	249	AR291: 6, AR296: 6, AR258: 6, AR274: 6, AR185: 6, AR218: 6, AR060: 6, AR293: 6, AR219: 5, AR256: 5, AR267: 5, AR259: 5, AR275: 5, AR269: 5, AR061: 5, AR240: 5, AR182: 5, AR237: 5, AR290: 5, AR249: 5, AR233: 5, AR273: 4, AR186: 4, AR104: 4, AR179: 3 H0265: 1, H0039: 1, H0616: 1, L0666: 1, L0748: 1 and H0506: 1.		
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AR281: 1, AR310: 0,							
AR104: 0							
S0126: 5, I 0766: 4							

64	HTEPC92	1091037	74	347 - 613	250	Gln-3 to Ser-29, Lys-56 to Asn-61, Asn-71 to Arg-77.	AR197: 6, AR241: 6, AR207: 5, AR162: 5, AR215: 5, AR163: 5		
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AR248:	1, AR204:	1,
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AR211:	1, AR203:	1,
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65	HAMFI04	1091042	75	682 - 1002	251	Gln-6 to Gly-12, Lys-27 to His-35, Arg-40 to Lys-49, Asn-71 to Asn-76, Asn-83 to Tyr-96, Pro-102 to Thr-107.	AR263: 6, AR184: 5, AR265: 5, AR251: 4, AR052: 4, AR198: 4, AR269: 4, AR309: 3, AR053: 3, AR270: 3, AR213: 3, AR268: 3, AR291: 3, AR267: 3, AR096: 3, AR183: 3, AR248: 3, AR290: 2, AR247: 2, AR312: 2, AR206: 2, AR289: 2, AR192: 2, AR204: 2, AR273: 2, AR266: 2, AR313: 2, AR182: 2, AR227: 1, AR229: 1,					

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AR185:	1, AR299:	1,
AR294:	1, AR232:	1,
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AR292:	1, AR284:	1,
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AR300:	1, AR061:	1,
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AR060:	1, AR226:	1,
AR243:	1, AR293:	1,
AR237:	1, AR256:	1,
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AR274:	1, AR259:	1,
AR218:	1, AR219:	1,
AR240:	1, AR233:	1,
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AR177:	0, AR246:	0,
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AR286:	0, AR275:	0,
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AR194:	0, AR314:	0
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66	HAIDK30	1091076	76	2 - 2167	252	Pro-52 to Thr-58, Pro-115 to Asp-122, Asp-190 to Gln-209.	L0748: 2, L0759: 2, S0114: 1, H0663: 1, L0717: 1, H0486: 1, H0421: 1, L0040: 1, H0252: 1, H0560: 1, L0763: 1, L0773: 1, S0374: 1, S0148: 1, H0520: 1, H0518: 1, H0436: 1, S0390: 1 and H0422: 1.		
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67	HCE1R41	1091358	77	21 - 446	253	<p>Glu-11 to Ile-16, Asn-18 to Tyr-23, His-64 to Thr-72, Gly-100 to Ala-120, Gln-122 to Gly-128, Pro-134 to Pro-142.</p>	<p>H0486: 1, H0090: 1, S0112: 1, T0042: 1, H0494: 1, S0210: 1, L0770: 1, L0662: 1, L0659: 1, L0665: 1, H0682: 1, H0445: 1, H0653: 1 and S0424: 1. AR204: 5, AR251: 4, AR186: 4, AR194: 4, AR274: 4, AR183: 3, AR253: 3, AR313: 3, AR060: 3, AR310: 3, AR055: 3, AR246: 3, AR269: 3, AR184: 3, AR185: 3, AR270: 3, AR052: 3, AR089: 3, AR182: 2, AR309: 2, AR266: 2, AR192: 2, AR265: 2, AR061: 2, AR282: 2, AR206: 2, AR291: 2, AR277: 2, AR312: 2, AR033: 2, AR104: 2, AR294: 2, AR284: 2, AR213: 2, AR283: 2, AR248: 2, AR296: 2, AR316: 2, AR298: 2, AR300: 2, AR039: 2, AR290: 2, AR286: 2, AR273: 2, AR292: 1, AR175: 1,</p>		
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68	HCESF90	1091570	78	4490 - 3606	254	Pro-74 to Pro-81, Arg-91 to Glu-96, Pro-139 to Asp-145, Ser-147 to His-157, Pro-167 to Leu-179, Pro-186 to Asp-197,	AR274: 15, AR213: 12, AR309: 12, AR229: 11, AR052: 11, AR183: 10, AR089: 10, AR313: 9, AR237: 8, AR271: 8, AR269: 8, AR244: 8,				

						Pro-204 to Ser-212, Val-247 to Phe-258, Pro-287 to Pro-292.	AR247: 7, AR238: 7, AR240: 7, AR194: 7, AR175: 7, AR192: 7, AR060: 7, AR053: 7, AR282: 7, AR310: 6, AR299: 6, AR226: 6, AR270: 6, AR312: 6, AR198: 6, AR249: 6, AR185: 5, AR290: 5, AR316: 5, AR265: 5, AR292: 5, AR182: 5, AR234: 5, AR263: 5, AR295: 5, AR184: 5, AR186: 5, AR177: 5, AR267: 4, AR055: 4, AR273: 4, AR205: 4, AR233: 4, AR096: 4, AR179: 4, AR061: 4, AR231: 4, AR268: 4, AR227: 4, AR104: 4, AR232: 4, AR293: 4, AR246: 4, AR033: 3, AR039: 3, AR300: 3, AR291: 3, AR277: 3, AR294: 3, AR289: 3, AR256: 3, AR296: 3, AR206: 3, AR253: 3, AR248: 3, AR298: 2, AR284: 2, AR286: 2, AR204: 2, AR275: 2,		
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69	HMSFT51	1091878	79	3 - 263	255	Pro-1 to Gln-8, Ser-17 to Gly-32, Ser-52 to Pro-60, Ser-77 to Gly-83.	S3014: 1, S0027: 1, L0748: 1, L0754: 1, L0749: 1, L0780: 1, L0589: 1, L0608: 1, H0667: 1, S0276: 1, H0543: 1, H0423: 1, S0458: 1, H0506: 1, UNKWN: 1 and H0352: 1.	7q33	180105, 222800
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70	HAICQ70	1091887	80	72 - 1508	256	His-10 to Ala-15, Arg-17 to Gln-22, Glu-69 to Thr-74, Asp-91 to Arg-99, Gln-122 to Cys-138, Ser-153 to Tyr-164,	AR261: 13, AR235: 12, AR311: 12, AR170: 12, AR168: 11, AR210: 11, AR308: 10, AR215: 10, AR214: 10, AR171: 10, AR161: 9, AR162: 9,			

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71	HDQZ89	1092002	81	75 - 581	257	Ser-9 to Gln-17, Pro-45 to Thr-53, Glu-93 to Glu-98.	L0794: 1, L0533: 1, L0379: 1, L0807: 1, L0512: 1, L0782: 1, L0384: 1, L0647: 1, L0788: 1, L0665: 1, S0374: 1, L0565: 1, L0438: 1, H0519: 1, H0689: 1, H0683: 1, H0684: 1, H0658: 1, S0330: 1, S0380: 1, L0602: 1, S0152: 1, S0013: 1, H0555: 1, H0631: 1, S0432: 1, S0037: 1, S0027: 1, S0028: 1, L0755: 1, L0757: 1, H0444: 1, L0596: 1, S0011: 1, S0026: 1, S0196: 1, S0456: 1 and H0008: 1. AR313: 6, AR263: 4, AR184: 4, AR312: 3, AR194: 3, AR253: 3, AR269: 3, AR096: 3, AR270: 3, AR052: 3, AR183: 3, AR310: 3, AR175: 3, AR309: 2, AR247: 2, AR282: 2, AR198: 2, AR296: 2, AR089: 2, AR192: 2, AR229: 2, AR268: 2,		
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72	HWAFW39	1092095	82	188 - 370	258			AR263: 8, AR183: 5, AR270: 5, AR182: 5, AR313: 4, AR184: 4, AR296: 4, AR269: 4, AR267: 3, AR268: 3, AR229: 3, AR285: 3, AR238: 3, AR290: 3, AR298: 3, AR202: 3, AR291: 3, AR310: 2, AR282: 2, AR251: 2, AR226: 2, AR089: 2, AR247: 2, AR286: 2, AR237: 2, AR266: 2, AR312: 2, AR231: 2, AR284: 2, AR175: 2, AR292: 2, AR246: 2, AR293: 2, AR277: 2, AR294: 2, AR060: 2, AR289: 2, AR299: 2, AR316: 2, AR096: 2, AR265: 2, AR234: 1, AR249: 1, AR240: 1,	5		

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73	HE8UL90	1092181	83	3 - 578	259				AR226: 6, AR170: 6, AR227: 4, AR238: 4, AR263: 4, AR232: 4, AR161: 3, AR061: 3, AR162: 3, AR163: 3, AR237: 3, AR168: 3, AR193: 3, AR166: 3,					

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74	HTEEF19	1092204	84	1521 - 1937	260	Ala-1 to Ser-7, Gln-108 to Ser-117.	AR275: 0, AR179: 0, AR164: 0, AR211: 0, AR173: 0, AR239: 0, AR241: 0, AR235: 0, AR178: 0, AR314: 0, AR297: 0, AR262: 0, AR230: 0, AR264: 0, AR203: 0 H0013: 1		
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77	HMERH77	1092363	87	2 - 844	263	Thr-4 to Ser-11	AR227: 2, AR284: 2, AR226: 2, AR287: 2, AR255: 2, AR234: 2, AR096: 2, AR168: 2, AR232: 2, AR277: 2, AR185: 1, AR191: 1, AR178: 1, AR283: 1, AR216: 1, AR285: 1, AR286: 1, AR262: 1, AR288: 1, AR186: 1, AR240: 1, AR292: 1, AR237: 1, AR231: 1, AR243: 1, AR204: 1, AR211: 1, AR215: 1, AR179: 1, AR190: 1, AR197: 1, AR188: 1, AR235: 1, AR104: 1, AR173: 1, AR290: 1, AR258: 1, AR192: 1, AR242: 1, AR294: 1, AR189: 1, AR296: 1, AR175: 1, AR260: 1, AR256: 1, AR202: 0, AR259: 0, AR250: 0, AR198: 0, AR249: 0, AR169: 0, AR174: 0, AR221: 0, AR244: 0, AR241: 0, AR206: 0 T0110: 1 and T0110: 1.		
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79	HLDAB75	1092493	89	1 - 1521	265	Gly-97 to Gly-108, Glu-175 to Val-180, Pro-189 to Gly-200, Arg-266 to Asp-272, Leu-295 to Asp-305, Glu-322 to Gly-330.	S0380: 1, S0152: 1, S0206: 1, L0786: 1, L0758: 1, H0595: 1, L0589: 1, L0603: 1, H0668: 1, H0665: 1, S0276: 1 and H0352: 1. AR055: 58, AR233: 52, AR226: 50, AR232: 49, AR192: 46, AR300: 45, AR237: 45, AR033: 44, AR204: 43, AR177: 43, AR294: 43, AR298: 42, AR061: 42, AR185: 41, AR186: 41, AR184: 40, AR284: 40, AR299: 40, AR198: 40, AR295: 39, AR267: 38, AR175: 37, AR234: 37, AR286: 37, AR293: 37, AR314: 36, AR285: 35, AR238: 33, AR182: 33, AR315: 33, AR206: 32, AR259: 31, AR205: 31, AR179: 31, AR227: 29, AR247: 29, AR282: 29, AR243: 28, AR283: 28, AR053: 27, AR104: 27, AR289: 27, AR291: 26, AR039: 26, AR275: 26, AR229: 26, AR258: 24, AR231: 24,	19cen- q13.2	
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81	HNTDM29	1092545	91	2 - 592	267	Pro-15 to Lys-27.	AR183: 6, AR265: 5, AR269: 5, AR270: 5, AR291: 4, AR296: 4, AR263: 4, AR282: 4, AR310: 4, AR268: 4, AR182: 3, AR266: 3, AR289: 3, AR313: 3, AR298: 3, AR294: 3,			

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82	HATCZ07	1093241	92	2 - 1111	268	Asn-1 to Val-6, Gly-50 to Phe-56, Val-74 to Asn-79	AR280: 1, AR184: 1, AR185: 1, AR314: 1, AR315: 1, AR246: 1, AR177: 1, AR243: 1, AR204: 1, AR179: 0, AR244: 0 S0222: 4, L0751: 4, L0769: 3, H0519: 3, H0052: 2, H0617: 2, H0658: 2, H0539: 2, L0750: 2, L0756: 2, H0624: 1, H0556: 1, S6024: 1, S0045: 1, S6026: 1, H0415: 1, H0592: 1, H0331: 1, H0156: 1, H0618: 1, H0620: 1, S0051: 1, S0038: 1, H0561: 1, L0803: 1, L0659: 1, L0809: 1, L0544: 1, L0787: 1, L0664: 1, L0665: 1, S0374: 1, L0438: 1, H0593: 1, H0435: 1, H0696: 1, L0777: 1, L0752: 1, L0366: 1, S0424: 1 and H0352: 1.	AR312: 36, AR269: 33, AR309: 33, AR052: 33, AR270: 27, AR291: 26	11q23.2	261640, 602574, 602574
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83	HDPXT07	1093488	93	87 - 1460	269	Ser-8 to Cys-18, Trp-52 to Gly-57,	AR271: 10, AR243: 10, AR253: 9, AR226: 9, AR248: 9, AR247: 9, AR256: 9, AR300: 9, AR227: 9, AR263: 8, AR055: 8, AR234: 8, AR231: 8, AR179: 8, AR233: 7, AR251: 6, AR229: 6, AR281: 0, AR280: 0 H0506: 4, H0619: 3, S0418: 2, H0156: 2, H0356: 2, H0038: 2, H0539: 2, L0759: 2, L0604: 2, S0040: 1, S0342: 1, H0295: 1, H0208: 1, H0486: 1, H0013: 1, H0599: 1, S0010: 1, H0052: 1, H0046: 1, H0050: 1, L0471: 1, H0024: 1, S0250: 1, H0031: 1, H0644: 1, H0591: 1, H0652: 1, L0532: 1, S0053: 1, S0126: 1, L0439: 1, L0756: 1, L0581: 1, L0603: 1 and S0026: 1.		
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84	HMCHG54	1093752	94	250 - 1407	270	Ser-1 to Arg-7, Thr-32 to Ser-43, Arg-163 to Gln-175, Lys-200 to Ser-206, Tyr-212 to Lys-219, Ser-254 to Tyr-266.	AR221: 6, AR225: 4, AR178: 3, AR254: 3, AR263: 3, AR172: 3, AR215: 2, AR060: 2, AR171: 2, AR257: 2, AR168: 2, AR161: 2, AR104: 2, AR192: 1, AR228: 1, AR096: 1, AR199: 1, AR261: 1, AR247: 1, AR162: 1, AR282: 1, AR274: 1, AR201: 1, AR165: 1,		

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AR242: 0, AR190: 0,							
AR232: 0							

85	HELGM30	1093937	95	2 - 661	271	His-1 to Asn-6, Arg-29 to Glu-34, Glu-152 to Asp-164, Gly-211 to Gly-216.	L0754: 6, L0766: 3, L0731: 2, H0624: 1, H0170: 1, S0116: 1, S0280: 1, H0545: 1, T0006: 1, S0344: 1, S0426: 1, L0770: 1, L0790: 1, L0748: 1, L0756: 1, L0779: 1, L0589: 1 and L0462: 1. AR192: 3, AR198: 2, AR241: 2, AR263: 2, AR284: 2, AR096: 2, AR282: 1, AR293: 1, AR312: 1, AR275: 1, AR183: 1, AR089: 1, AR194: 1, AR269: 1, AR247: 1, AR277: 1, AR177: 1, AR296: 1, AR060: 1, AR227: 1, AR244: 1, AR310: 1, AR300: 1, AR055: 1, AR104: 1, AR186: 1, AR294: 1, AR316: 1, AR299: 1, AR175: 0, AR233: 0, AR248: 0, AR185: 0, AR313: 0, AR061: 0, AR280: 0, AR053: 0, AR274: 0, AR243: 0, AR249: 0, AR246: 0, AR251: 0.		
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[illegible]

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86	HLKAB61	1094201	96	79 - 453	272	Gln-28 to Tyr-33, His-46 to Tyr-51, Tyr-62 to Gly-78, Gly-84 to Asp-91.	AR280: 8, AR315: 8, AR183: 8, AR314: 7, AR281: 7, AR270: 6, AR263: 6, AR184: 5, AR291: 5, AR268: 5,	1p13-p11	102770, 115200, 132810, 132810, 188540,
							H0231: 1, H0150: 1, H0019: 1, L0471: 1, H0051: 1, H0071: 1, H0416: 1, H0292: 1, H0252: 1, L0483: 1, H0213: 1, H0040: 1, H0551: 1, H0413: 1, L0564: 1, S0002: 1, L0763: 1, L0770: 1, L0761: 1, L0646: 1, L0800: 1, L0643: 1, L0771: 1, L0651: 1, L0653: 1, L0655: 1, L0658: 1, L0518: 1, L0383: 1, L0809: 1, L0787: 1, L0666: 1, L0664: 1, L0352: 1, H0519: 1, H0648: 1, H0518: 1, S0152: 1, H0521: 1, S0190: 1, S0392: 1, S3014: 1, L0744: 1, L0754: 1, L0758: 1, L0759: 1, L0592: 1, S0192: 1 and H0543: 1.		

AR269:	5, AR229:	4,	600234,
AR247:	4, AR175:	4,	601414,
AR292:	4, AR290:	4,	601691,
AR231:	4, AR182:	4,	601691,
AR226:	4, AR313:	4,	601691,
AR177:	4, AR295:	4,	601691,
AR089:	4, AR234:	4,	601718,
AR238:	4, AR246:	3,	602094
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AR298:	3, AR253:	3,	
AR294:	3, AR285:	3,	
AR265:	3, AR277:	3,	
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AR299:	2, AR289:	2,	
AR286:	2, AR219:	2,	
AR248:	2, AR284:	2,	
AR232:	2, AR296:	2,	
AR053:	2, AR266:	2,	
AR240:	2, AR249:	2,	
AR186:	2, AR218:	2,	
AR309:	2, AR033:	2,	
AR259:	2, AR233:	2,	
AR310:	2, AR052:	2,	
AR213:	1, AR061:	1,	
AR312:	1, AR060:	1,	
AR256:	1, AR243:	1,	
AR275:	1, AR192:	1,	
AR300:	1, AR185:	1,	

87	HFIZB56	1094512	97	3 - 845	273	Thr-23 to Ser-30, Arg-58 to Asp-64, Ala-75 to Asn-82, Glu-103 to Gln-112, Leu-119 to Thr-141, Lys-175 to Tyr-182, Asn-211 to Thr-217.	AR055: 1, AR258: 1, AR241: 1, AR039: 1, AR283: 1, AR194: 1, AR205: 1, AR251: 1, AR198: 0, AR244: 0, AR204: 0, AR273: 0, AR202: 0, AR206: 0, AR274: 0, AR104: 0, AR271: 0 H0386: 2 and H0610: 1. AR241: 41, AR313: 25, AR194: 20, AR192: 18, AR229: 18, AR173: 18, AR198: 17, AR258: 17, AR269: 17, AR183: 17, AR039: 15, AR244: 15, AR089: 15, AR202: 15, AR219: 15, AR182: 14, AR184: 14, AR218: 14, AR161: 13, AR162: 13, AR265: 13, AR240: 13, AR096: 13, AR296: 13, AR293: 13, AR163: 12, AR238: 12, AR165: 12, AR053: 12, AR292: 12, AR204: 12, AR299: 12, AR175: 12, AR285: 12, AR164: 12, AR257: 12, AR236: 12, AR262: 12, AR247: 11, AR206: 11,	9q31-q32	109400, 132800, 132800, 154400, 186855, 223900, 253800, 253800, 278700, 602088
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88	HAGFJ46	1094598	98	2 - 1330	274	Pro-273 to Cys-278,	AR261: 6, AR248: 6, AR203: 6, AR060: 6, AR255: 6, AR230: 6, AR171: 6, AR225: 6, AR215: 6, AR266: 6, AR256: 6, AR176: 5, AR242: 5, AR288: 5, AR207: 5, AR188: 5, AR221: 5, AR235: 5, AR170: 5, AR252: 5, AR172: 5, AR309: 5, AR214: 5, AR228: 5, AR224: 5, AR212: 5, AR197: 5, AR232: 4, AR223: 4, AR311: 4, AR190: 4, AR222: 4, AR253: 4, AR193: 4, AR211: 4, AR239: 4, AR283: 4, AR308: 4, AR216: 4, AR168: 4, AR217: 4, AR210: 3, AR195: 3, AR272: 3, AR061: 2, AR169: 2, AR055: 2, AR245: 2, AR201: 2, AR254: 1, AR250: 0 S0250: 1, H0030: 1, H0521: 1, S0192: 1 and S0242: 1.	7q31	126650,
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Leu-399 to Ser-405, Glu-416 to Pro-430, Ser-433 to Thr-441.	AR193:1014, AR181: 924, AR178: 868, AR255: 867, AR201: 863, AR199: 787, AR228: 779, AR191: 756, AR287: 747, AR262: 745, AR177: 726, AR257: 723, AR179: 700, AR261: 673, AR288: 664, AR197: 663, AR233: 663, AR174: 593, AR196: 587, AR260: 586, AR230: 578, AR190: 569, AR188: 568, AR175: 554, AR242: 537, AR286: 535, AR294: 533, AR296: 526, AR176: 524, AR203: 513, AR235: 508, AR173: 496, AR055: 492, AR060: 482, AR258: 476, AR299: 472, AR195: 469, AR189: 448, AR291: 439, AR284: 439, AR226: 436, AR300: 431, AR231: 424, AR297: 421 AR033: 420	126650, 164860, 180105, 222800, 246900, 274600, 274600, 602081
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89	HYAAM43	1094663	99	2 - 805	275	Glu-1 to Gln-6, Glu-17 to Phe-66, Thr-100 to Phe-106, Cys-117 to Ser-124, Asn-132 to Ser-138, Ser-169 to Glu-174, Lys-179 to Arg-184, Arg-232 to Asp-243, Pro-261 to Ser-267.	AR251: 87, AR194: 61, AR248: 58, AR281: 54, AR253: 54, AR202: 48, AR314: 41, AR315: 40, AR039: 40, AR205: 37, AR246: 36, AR206: 35, AR244: 34, AR280: 33, AR263: 28, AR283: 27, AR241: 26, AR243: 26, AR192: 25, AR271: 24, AR265: 24, AR033: 24, AR096: 22, AR198: 22,		

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90	HDAAN31	1094668	100	32 - 718	276	Arg-11 to Val-19, Ser-76 to Pro-81, Thr-121 to Thr-128, Gln-163 to Gly-169, Glu-179 to Thr-186, Lys-191 to Lys-197, Ala-202 to Val-207.	AR290: 6, AR179: 5, AR267: 5 L0794: 5, L0804: 3, H0250: 2, H0271: 2, S0002: 2, S0052: 2, H0583: 1, H0427: 1, H0042: 1, H0421: 1, H0264: 1, L0637: 1, L0800: 1, L0790: 1, L0791: 1, S0216: 1 and H0555: 1.		
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91	HNGEO79	1094775	101	200 - 709	277	Gln-41 to Asp-48, Ser-67 to Leu-72, Thr-79 to Ser-84, Ile-96 to Ser-103, Arg-120 to Ser-127.	S0282: 1, H0497: 1, H0616: 1, L0807: 1, L0438: 1, H0520: 1, H0539: 1, L0740: 1, L0779: 1, L0777: 1 and L0589: 1. AR249: 40, AR186: 40, AR251: 30, AR061: 29, AR241: 27, AR229: 27, AR033: 27, AR292: 26, AR206: 25, AR283: 25, AR294: 24, AR204: 22, AR055: 21, AR237: 21, AR185: 21, AR233: 21, AR104: 21, AR293: 19, AR248: 19, AR282: 19, AR227: 19, AR275: 18, AR273: 18, AR052: 18, AR300: 17, AR271: 17, AR192: 17, AR194: 17, AR314: 16, AR198: 16, AR274: 16, AR089: 16, AR184: 15, AR231: 15, AR243: 15, AR232: 14, AR244: 14, AR060: 14, AR313: 14, AR316: 14, AR234: 13, AR183: 13, AR096: 13, AR281: 13, AR315: 12, AR253: 12, AR280: 12, AR238: 12,		
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92	HWHQR25	1094777	102	2 - 502	278	Asn-7 to Thr-16, Cys-20 to Trp-26, Gln-33 to Asn-46, Ala-48 to Tyr-58, Glu-77 to Tyr-96, Gly-125 to Asn-135.	AR221: 3, AR164: 3, AR288: 3, AR171: 3, AR193: 3, AR250: 3, AR239: 3, AR166: 3, AR230: 3, AR311: 3, AR254: 3, AR199: 3, AR222: 3, AR262: 3, AR297: 2, AR245: 2, AR173: 2, AR224: 2, AR195: 2, AR258: 2, AR216: 2, AR255: 2, AR174: 2, AR196: 2, AR191: 2, AR189: 2, AR200: 2, AR172: 2, AR219: 2, AR218: 2, AR217: 2, AR212: 2, AR190: 2, AR188: 1, AR308: 1, AR203: 1, AR260: 1, AR242: 1, AR211: 1, AR168: 1, AR252: 1, AR210: 0, AR170: 0 S0052: 1 and L0758: 1.	15q14	102540, 118511, 218000, 227220, 243500, 254770, 601800
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AR188:	1, AR232:	1,
AR289:	1, AR268:	1,
AR186:	1, AR266:	1,
AR298:	1, AR291:	1,
AR210:	1, AR285:	1,
AR295:	1, AR172:	1,
AR237:	1, AR294:	1,
AR251:	1, AR290:	1,
AR297:	1, AR238:	1,
AR296:	1, AR175:	1,
AR267:	1, AR174:	1,
AR229:	1, AR257:	1,
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AR177:	1, AR292:	1,
AR260:	1, AR293:	1,
AR281:	1, AR224:	1,
AR194:	1, AR215:	0,
AR231:	0, AR217:	0,
AR261:	0, AR248:	0,
AR176:	0, AR314:	0,
AR233:	0, AR202:	0,
AR219:	0, AR218:	0,
AR236:	0, AR256:	0,
AR225:	0, AR206:	0,
AR288:	0, AR223:	0,
AR181:	0, AR173:	0,
AR258:	0, AR280:	0,
AR273:	0, AR239:	0,
AR222:	0, AR255:	0,

93	HNTAV78	1094783	103	3 - 266	279	Glu-52 to Leu-58, Arg-63 to Lys-71, Arg-83 to Val-88.	AR178: 0, AR228: 0, AR199: 0, AR259: 0, AR226: 0, AR196: 0, AR052: 0, AR234: 0, AR190: 0, AR191: 0, AR315: 0, AR189: 0 L0794: 5, L0777: 4, H0509: 2, L0772: 2, L0804: 2, H0144: 2, L0754: 2, L0747: 2, S0045: 1, S6022: 1, H0392: 1, H0592: 1, H0587: 1, H0486: 1, T0010: 1, H0623: 1, L0800: 1, L0643: 1, L0764: 1, L0768: 1, L0803: 1, L0653: 1, L0665: 1 and L0750: 1.	13q14.12- 21.1	180200, 180200, 180200, 180200, 256731, 277900, 600631, 602085
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AR293:	2, AR214:	2,						
AR182:	1, AR175:	1,						
AR294:	1, AR229:	1,						
AR300:	1, AR226:	1,						

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94	HNTCU89	1094785	104	2 - 550	280	Ile-26 to Gly-31, Arg-161 to Thr-168.	AR188: 0 H0305: 1, H0580: 1, H0428: 1, L0803: 1, L0809: 1 and H0519: 1. AR194: 13, AR263: 11, AR205: 10, AR246: 9, AR202: 9, AR206: 9, AR244: 9, AR039: 8, AR281: 7, AR315: 6, AR243: 6, AR241: 6, AR204: 6, AR275: 6, AR282: 5, AR192: 5, AR175: 5, AR280: 5, AR271: 4, AR198: 4, AR310: 4, AR240: 4, AR295: 4, AR270: 4, AR265: 4, AR314: 4, AR053: 4, AR251: 4, AR273: 4, AR283: 4, AR269: 3, AR177: 3, AR213: 3, AR299: 3, AR316: 3, AR291: 3, AR285: 3, AR277: 3, AR104: 3, AR247: 3, AR182: 3, AR033: 3, AR266: 3, AR292: 3, AR096: 3, AR284: 3, AR183: 3, AR289: 3, AR312: 3, AR232: 3, AR052: 3, AR268: 3,	1		
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95	HTSHF07	1094859	105	1 - 762	281	Ala-53 to Ala-63,	H0436: 2, L0756: 2, L0362: 2, H0543: 2, H0624: 1, L0002: 1, H0650: 1, S0418: 1, H0208: 1, S0045: 1, S0046: 1, H0619: 1, T0048: 1, H0318: 1, L0109: 1, H0046: 1, H0050: 1, L0471: 1, S0051: 1, H0083: 1, H0328: 1, H0604: 1, H0068: 1, H0038: 1, H0551: 1, H0560: 1, H0561: 1, H0132: 1, L0763: 1, L0667: 1, L0662: 1, L0388: 1, L0804: 1, L0805: 1, L0606: 1, L0657: 1, L0659: 1, L0542: 1, L0518: 1, L0809: 1, L0791: 1, L0792: 1, H0547: 1, H0519: 1, H0682: 1, H0648: 1, S0330: 1, H0696: 1, L0750: 1, L0779: 1, L0731: 1, L0757: 1, L0759: 1, H0445: 1, L0480: 1, H0542: 1 and H0506: 1.	AR221: 9, AR176: 8,	11q13	102200,
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Pro-66 to Cys-71, Pro-73 to His-82, Pro-89 to Glu-102, Asp-158 to Glu-167, Arg-199 to Cys-231.	AR172: 7, AR268: 7, AR290: 7, AR222: 6, AR269: 6, AR183: 6, AR265: 5, AR182: 5, AR224: 5, AR211: 5, AR171: 5, AR178: 5, AR210: 4, AR310: 4, AR180: 4, AR267: 4, AR189: 4, AR175: 4, AR177: 4, AR270: 4, AR217: 4, AR181: 4, AR162: 4, AR161: 4, AR163: 4, AR247: 3, AR173: 3, AR216: 3, AR184: 3, AR245: 3, AR264: 3, AR289: 3, AR257: 3, AR311: 3, AR225: 3, AR223: 3, AR288: 3, AR165: 3, AR296: 2, AR266: 2, AR052: 2, AR292: 2, AR199: 2, AR089: 2, AR282: 2, AR200: 2, AR166: 2, AR249: 2, AR228: 2, AR061: 2, AR236: 2, AR230: 2, AR191: 2, AR284: 2, AR190: 2, AR293: 2, AR174: 2, AR291: 2, AR263: 2, AR252: 2,	106100, 131100, 131100, 131100, 133780, 147050, 153700, 161015, 164009, 168461, 168461, 168461, 180721, 180840, 191181, 193235, 209901, 232600, 259700, 259770, 600045, 600319, 600528, 601884
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AR213:	1, AR234:	1,
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96	HISAF60	1094890	106	4001 - 4261	282	Pro-8 to Asn-18, Gln-25 to Glu-30, Arg-50 to Arg-65.	L0659: 1, L0384: 1, L0809: 1, L0565: 1, H0547: 1, H0658: 1, S0037: 1, S0028: 1, S0206: 1, L0603: 1 and H0665: 1. AR194: 15, AR243: 14, AR033: 14, AR039: 13, AR274: 13, AR202: 12, AR246: 11, AR104: 11, AR263: 11, AR206: 11, AR205: 11, AR270: 10, AR265: 10, AR313: 10, AR053: 10, AR281: 10, AR198: 10, AR241: 10, AR299: 10, AR192: 10, AR247: 10, AR204: 9, AR244: 9, AR089: 9, AR271: 9, AR213: 9, AR312: 9, AR269: 9, AR052: 9, AR310: 9, AR284: 9, AR096: 8, AR275: 8, AR300: 8, AR285: 8, AR289: 8, AR315: 8, AR314: 8, AR309: 8, AR184: 7, AR240: 7, AR283: 7, AR266: 7, AR316: 7, AR280: 7, AR268: 7, AR182: 7, AR186: 7,			
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97	HWAGT47	1094891	107	63 - 608	283				L0588: 2, H0716: 1, L0002: 1, L0443: 1, S0001: 1, S0360: 1, H0645: 1, H0411: 1, H0587: 1, H0333: 1, H0486: 1, S0010: 1, S0050: 1, H0051: 1, H0428: 1, H0553: 1, H0032: 1, L0455: 1, S0036: 1, H0038: 1, H0412: 1, H0413: 1, H0100: 1, T0042: 1, L0770: 1, L0637: 1, L0766: 1, L0649: 1, L0774: 1, L0776: 1, L0655: 1, L0659: 1, L0783: 1, L0529: 1, L5623: 1, L0790: 1, L0791: 1, H0520: 1, H0519: 1, H0593: 1, H0689: 1, H0670: 1, H0672: 1, H0696: 1, S3014: 1, L0741: 1, L0744: 1, L0757: 1, L0608: 1 and S0398: 1.	12p12.3- 13.2	103950, 200990, 601458, 602096
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98	HSXDF41	1094953	108	3 - 1655	284	Arg-16 to Gly-23, Cys-30 to Ala-36, Asp-52 to Asp-58, Thr-98 to Gly-103			AR104: 24, AR033: 22, AR251: 20, AR186: 17, AR271: 13, AR290: 13, AR273: 12, AR182: 12	11				

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99	HMTMD88	1095222	109	1 - 750	285	Arg-6 to Gly-14, Gln-52 to Cys-58, Phe-93 to Gly-107, Ser-116 to Asn-121, Pro-134 to Gln-141, Ser-150 to Gly-165, Arg-197 to Trp-204, Asn-213 to Asn-222, Arg-243 to Lys-250.	AR286: 5, AR177: 5, AR315: 5, AR259: 5, AR294: 5, AR234: 5, AR061: 5, AR194: 5, AR233: 4, AR229: 4, AR241: 4, AR277: 4, AR258: 4, AR247: 4, AR256: 3, AR283: 3, AR179: 3, AR227: 2, AR281: 1		
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100	HENEA55	1095289	110	2 - 766	286	Lys-14 to Ser-20.	AR221: 56, AR225: 55, AR223: 51, AR215: 48, AR217: 46, AR171: 36, AR216: 31, AR224: 29, AR170: 28, AR222: 27, AR169: 25, AR214: 24, AR168: 24, AR172: 23, AR183: 16, AR269: 15, AR270: 15, AR268: 14, AR173: 13, AR180: 13, AR219: 12, AR182: 12, AR290: 12, AR161: 11, AR162: 11, AR163: 11, AR165: 10, AR164: 9, AR166: 9, AR181: 9, AR178: 9, AR291: 9, AR184: 8, AR175: 8, AR297: 8, AR255: 7, AR089: 7, AR296: 7, AR292: 7, AR210: 7, AR287: 6, AR191: 6, AR179: 6, AR190: 6, AR316: 6, AR250: 6, AR267: 6, AR285: 6, AR257: 6, AR312: 6, AR298: 6, AR248: 6, AR176: 5, AR293: 5, AR300: 5, AR254: 5, AR218: 5, AR313: 5, AR308: 5, AR294: 5,		
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101	HLJBH31	1095303	111	20 - 2146	287		H0659: 1, H0672: 1, S3014: 1, S0028: 1, L0752: 1, H0595: 1, S0394: 1, L0596: 1, L0589: 1, L0485: 1, L0595: 1, H0667: 1, S0242: 1, S0194: 1, H0423: 1, S0384: 1, H0506: 1 and H0352: 1.		
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102	HFTCV04	1095439	112	277 - 1050	288	Asp-35 to Ser-56, Pro-59 to His-66, Gln-111 to Lys-124	AR247: 5, AR055: 5, AR228: 5, AR293: 5, AR232: 5, AR275: 5, AR280: 5, AR253: 5, AR294: 5, AR314: 5, AR290: 5, AR218: 5, AR219: 5, AR183: 5, AR226: 5, AR244: 5, AR227: 5, AR251: 5, AR234: 5, AR231: 5, AR237: 5, AR270: 4, AR243: 4, AR204: 4, AR268: 4, AR265: 4, AR266: 4, AR274: 4, AR310: 4, AR233: 4, AR256: 4, AR175: 4, AR269: 4, AR206: 4, AR179: 3, AR267: 3, AR182: 3, AR061: 3, AR052: 3, AR186: 2, AR248: 2, AR273: 2, AR292: 2, AR284: 1, AR298: 1, AR259: 1, AR249: 1, AR184: 1, AR241: 1 H0375: 1, H0625: 1, H0529: 1 and S0216: 1.	AR251: 14, AR273: 13, 9q22 AR052: 9, AR310: 9, AR241: 9, AR265: 8,	229600, 264300, 600542,
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AR280:	0, AR261:	0,						
AR207:	0, AR169:	0,						
AR176:	0, AR260:	0,						

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103	HMCHH31	1095550	113	85 - 1215	289	<p>Thr-66 to Arg-74, Pro-140 to Glu-145, Lys-153 to Cys-161, Ala-198 to Gly-203, Ala-248 to Tyr-256, Thr-262 to Gln-273, Pro-292 to Ser-297, Thr-302 to Phe-324.</p>	<p>L0598: 1, L0763: 1, L0769: 1, L0521: 1, L0662: 1, L0558: 1, H0658: 1, S0380: 1, S0332: 1, S0028: 1, L0779: 1, L0758: 1, H0444: 1, S0192: 1, S0196: 1 and S0458: 1, AR213: 10, AR052: 7, AR192: 6, AR204: 6, AR251: 6, AR202: 6, AR053: 5, AR274: 5, AR246: 5, AR186: 5, AR055: 5, AR312: 5, AR184: 5, AR244: 5, AR060: 5, AR273: 5, AR061: 5, AR243: 5, AR270: 5, AR313: 4, AR206: 4, AR309: 4, AR185: 4, AR269: 4, AR267: 4, AR253: 4, AR275: 3, AR290: 3, AR033: 3, AR182: 3, AR096: 3, AR198: 3, AR194: 3, AR282: 3, AR316: 3, AR249: 3, AR291: 3, AR310: 3, AR295: 3, AR039: 3, AR296: 3, AR089: 3, AR283: 3, AR183: 3,</p>		
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104	HNECM88	1095640	114	1 - 753	290	Leu-9 to Arg-16, Asp-44 to Ser-51, Pro-57 to Thr-64, Asp-132 to Ala-137, Leu-148 to Met-155, Arg-238 to Gly-251.	L0794: 1, L0789: 1, S0053: 1, H0689: 1, H0134: 1, L0779: 1, L0777: 1 and H0445: 1. AR248: 10, AR183: 9, AR313: 8, AR033: 7, AR253: 7, AR241: 7, AR096: 7, AR275: 6, AR281: 6, AR263: 6, AR186: 6, AR192: 6, AR089: 6, AR218: 6, AR104: 5, AR249: 5, AR258: 5, AR238: 5, AR310: 5, AR247: 5, AR299: 5, AR312: 5, AR293: 5, AR175: 5, AR185: 5, AR229: 5, AR274: 5, AR055: 5, AR240: 5, AR314: 4, AR316: 4, AR202: 4, AR219: 4, AR300: 4, AR292: 4, AR060: 4, AR282: 4, AR265: 4, AR061: 4, AR213: 4, AR243: 4, AR315: 4, AR234: 4, AR053: 4, AR280: 4, AR259: 4, AR246: 4, AR271: 4, AR273: 4, AR277: 4, AR283: 4, AR194: 4,	19q13.3	113900, 126340, 126391, 130410, 134790, 138570, 160900, 173850, 258501, 600040, 602225, 602225
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105	HCQBI77	1095827	115	983 - 267	291				H0038: 1, H0412: 1, H0059: 1, L0762: 1, L0770: 1, L0796: 1, L0761: 1, L0764: 1, L0662: 1, L0767: 1, L0649: 1, L0804: 1, L0650: 1, L0806: 1, L0606: 1, L0783: 1, L0809: 1, L0665: 1, S0053: 1, H0659: 1, S0330: 1, L0602: 1, H0694: 1, S3014: 1, S0028: 1, L0740: 1, L0754: 1, L0756: 1, L0752: 1 and L0588: 1.		
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106	HBXFI23	1095877	116	196 - 549	292	Gly-6 to Pro-19, Arg-67 to Tyr-74.	L0779: 3, H0087: 2, L0770: 2, L0375: 2, L0659: 2, L0747: 2, L0601: 2, H0556: 1, H0657: 1, H0253: 1, H0263: 1, H0545: 1, L0471: 1, H0594: 1, T0041: 1, L0451: 1, L0369: 1, L0762: 1, L0662: 1, L0499: 1, L0651: 1, L0806: 1, L0776: 1, L0513: 1, L0809: 1, L0791: 1, L0663: 1, L0664: 1, L0438: 1, L0741: 1, L0748: 1, L0740: 1, L0756: 1, L0780: 1, L0755: 1, L0759: 1 and H0542: 1.		
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AR262:	2, AR180:	2,
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AR211:	1, AR253:	1,
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AR193:	1, AR055:	1,
AR284:	1, AR195:	1,
AR258:	1, AR275:	1,
AR308:	1, AR246:	1,
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AR241:	1, AR244:	1,
AR219:	0, AR198:	0,
AR256:	0, AR281:	0,
AR194:	0, AR243:	0,
AR254:	0, AR242:	0,
AR039:	0, AR314:	0,
AR170:	0, AR190:	0,
AR197:	0, AR245:	0,
AR218:	0, AR315:	0,

107	HEOAD12	1095888	117	332 - 934	293	Pro-35 to Trp-40, Pro-54 to Pro-59, Lys-91 to Met-122.	AR207: 0, AR199: 0, AR251: 0, AR280: 0, AR250: 0, AR259: 0, AR201: 0, AR235: 0 L0748: 3, L0752: 3, L0740: 2, L0731: 2, S0358: 1, H0438: 1, H0574: 1, H0421: 1, H0046: 1, H0041: 1, H0063: 1, H0272: 1, S0150: 1, S0002: 1, L0794: 1, L0803: 1, L0804: 1, L0775: 1, L0661: 1, L0789: 1, H0672: 1, H0539: 1, L0758: 1 and L0366: 1.	4q28-q32	107250, 134820, 134820, 134820, 134830, 134850, 134850, 181600, 189800, 208400, 231675, 266300
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AR216:	2, AR277:	2,						
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AR242: 1, AR207: 1,									
AR170: 1, AR248: 1,									
AR210: 1, AR281: 0,									
AR272: 0, AR260: 0,									
AR230: 0									

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108	HEMFC27	1095935	118	69 - 1100	294	Ile-122 to Lys-130, Tyr-151 to Thr-158, Gln-176 to Thr-181, Ile-215 to Lys-220, Thr-233 to Thr-238, Thr-291 to Glu-298, Val-304 to Thr-335.	H0547: 1, H0659: 1, - S0380: 1, H0710: 1, S0190: 1, H0727: 1, L0754: 1, L0756: 1, L0592: 1, H0667: 1, H0136: 1 and S0276: 1. AR268: 36, AR290: 29, AR269: 18, AR270: 18, AR267: 16, AR182: 11, AR238: 10, AR184: 9, AR096: 8, AR175: 7, AR192: 7, AR295: 7, AR292: 7, AR183: 7, AR227: 6, AR291: 6, AR244: 6, AR299: 6, AR246: 5, AR185: 5, AR275: 5, AR284: 5, AR186: 5, AR060: 4, AR053: 4, AR273: 4, AR316: 4, AR266: 4, AR253: 4, AR313: 4, AR247: 4, AR289: 4, AR240: 4, AR293: 4, AR177: 4, AR061: 4, AR282: 3, AR194: 3, AR055: 3, AR052: 3, AR179: 3, AR309: 3, AR285: 3, AR204: 3, AR271: 3, AR089: 3, AR033: 3, AR213: 3,	11p15	108985, 186921, 602092
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109	HFOYM21	1096023	119	1 - 3174	295	Pro-10 to Asp-17, Phe-47 to Arg-55, Leu-86 to Asp-96, Val-120 to Asp-125, Thr-135 to Asn-140, Glu-152 to Thr-159, Val-166 to Ser-173, Asn-241 to Val-246, Glu-257 to Arg-265, Gly-296 to Leu-304, Gln-310 to Thr-317, Tyr-327 to Ile-332, Gly-378 to Thr-384, Glu-432 to Lys-443, Glu-532 to Lys-538, Arg-545 to Val-550,	S0360: 1, S0046: 1, H0351: 1, H0411: 1, H0013: 1, H0244: 1, L0105: 1, H0172: 1, H0123: 1, H0050: 1, H0355: 1, H0375: 1, H0328: 1, H0644: 1, H0212: 1, H0124: 1, H0616: 1, H0412: 1, H0413: 1, L0371: 1, L0790: 1, H0519: 1, L0779: 1, L0755: 1, L0759: 1, S0260: 1, H0595: 1 and H0506: 1, AR263: 13, AR184: 7, AR215: 6, AR194: 6, AR265: 5, AR224: 5, AR180: 5, AR171: 5, AR183: 4, AR310: 4, AR214: 4, AR168: 4, AR216: 4, AR197: 4, AR264: 4, AR207: 4, AR223: 3, AR308: 3, AR289: 3, AR277: 3, AR170: 3, AR202: 3, AR282: 3, AR235: 3, AR052: 3, AR312: 3, AR238: 3, AR206: 3, AR165: 3, AR266: 3, AR311: 3, AR228: 3,	15q22.3- q23	118485, 151670, 231680, 272800, 272800, 272800, 276700, 600374, 601780
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				Gly-553 to Phe-566, Asp-575 to Tyr-581, Thr-611 to Thr-617, His-680 to Asp-689, Met-721 to Arg-729, Asp-733 to Thr-740, Glu-779 to Gly-784, Asn-803 to His-810, Leu-834 to Asp-840, Leu-868 to Ala-875, Gln-895 to Gly-901, Glu-908 to Gln-913, Ser-955 to Asn-967, Ser-998 to Ile-1006.	AR164: 3, AR295: 3, AR296: 3, AR166: 3, AR225: 3, AR275: 3, AR251: 3, AR254: 3, AR288: 2, AR222: 2, AR261: 2, AR198: 2, AR270: 2, AR217: 2, AR297: 2, AR247: 2, AR176: 2, AR053: 2, AR299: 2, AR257: 2, AR196: 2, AR239: 2, AR316: 2, AR269: 2, AR244: 2, AR163: 2, AR309: 2, AR286: 2, AR089: 2, AR055: 2, AR162: 2, AR169: 2, AR234: 2, AR283: 2, AR237: 2, AR096: 2, AR192: 2, AR221: 2, AR229: 2, AR268: 2, AR290: 2, AR281: 2, AR186: 2, AR174: 2, AR182: 2, AR230: 2, AR033: 2, AR188: 2, AR313: 2, AR246: 2, AR173: 2, AR292: 2, AR200: 2, AR240: 2, AR213: 2, AR190: 2, AR285: 2, AR212: 2, AR231: 2, AR272: 2,			
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AR256:	1, AR255:	1,							
AR203:	1, AR039:	1,							
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AR271:	1, AR249:	1,							
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AR193:	0	0							

110	HCE3H71	1096278	120	2 - 823	296	Pro-39 to Glu-46, Pro-51 to Gln-59.	S0126: 8, S0250: 7, H0251: 3, H0545: 2, L0794: 2, L0565: 2, L0744: 2, L0757: 2, S0040: 1, S0212: 1, S0418: 1, S0360: 1, H0549: 1, H0530: 1, H0024: 1, H0252: 1, L0053: 1, H0124: 1, S0208: 1 and S0276: 1, AR202: 63, AR194: 63, AR205: 59, AR039: 53, AR206: 51, AR204: 48, AR192: 46, AR244: 46, AR243: 46, AR246: 44, AR198: 42, AR241: 39, AR271: 30, AR053: 29, AR273: 29, AR274: 26, AR265: 26, AR213: 25, AR310: 24, AR263: 23, AR282: 22, AR275: 22, AR104: 21, AR309: 21, AR240: 19, AR312: 19, AR295: 19, AR052: 19, AR251: 17, AR316: 17, AR313: 16, AR055: 16, AR186: 16, AR033: 15, AR096: 15, AR285: 15, AR060: 15, AR299: 14, AR284: 14, AR277: 14,		
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111	HOABR66	1096304	121	3 - 440	297	Ser-1 to Glu-7, Gln-56 to Trp-66, Thr-72 to Phe-79.	L0786: 1. AR274: 9, AR194: 8, AR244: 8, AR268: 8, AR183: 8, AR275: 8, AR198: 8, AR269: 7, AR290: 7, AR281: 7, AR270: 7, AR206: 7, AR186: 7, AR192: 7, AR055: 6, AR265: 6, AR273: 6, AR182: 6, AR247: 6, AR184: 6, AR060: 5, AR204: 5, AR052: 5, AR175: 5, AR053: 5, AR313: 5, AR240: 5, AR310: 5, AR267: 5, AR177: 5, AR263: 5, AR246: 5, AR089: 5, AR061: 5, AR280: 4, AR282: 4, AR315: 4, AR316: 4, AR243: 4, AR289: 4, AR314: 4, AR271: 4, AR277: 4, AR299: 4, AR202: 4, AR033: 4, AR286: 4, AR300: 4, AR205: 4, AR185: 4, AR312: 4, AR266: 4, AR241: 3, AR298: 3, AR104: 3, AR096: 3, AR213: 3, AR291: 3,	17	
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AR270:	3, AR265:	3,
AR282:	2, AR286:	2,
AR298:	2, AR266:	2,
AR267:	1, AR290:	1,
AR227:	1, AR309:	1,
AR247:	1, AR096:	1,
AR294:	1, AR312:	1,
AR269:	1, AR089:	1,
AR055:	1, AR232:	1,
AR284:	1, AR231:	1,
AR316:	1, AR198:	1,
AR283:	1, AR300:	0,
AR299:	0, AR248:	0,
AR033:	0, AR249:	0,
AR060:	0, AR285:	0,
AR185:	0, AR177:	0,
AR182:	0, AR291:	0,
AR243:	0, AR192:	0,
AR295:	0, AR277:	0,
AR273:	0, AR314:	0,
AR218:	0, AR206:	0,
AR256:	0, AR237:	0,
AR229:	0, AR274:	0,
AR289:	0, AR186:	0,
AR052:	0, AR204:	0,
AR280:	0, AR205:	0,
AR202:	0, AR268:	0,
AR233:	0, AR315:	0,
AR313:	0, AR104:	0,
AR234:	0, AR179:	0,

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113	HDPKI84	1096306	123	62 - 802	299	Gln-1 to His-8, Ile-65 to Trp-71, Pro-101 to Glu-109, Ser-149 to Lys-154, Lys-181 to Gly-188.	AR227: 26, AR263: 16, AR232: 15, AR298: 15, AR284: 14, AR182: 14, AR229: 13, AR266: 10, AR238: 7, AR289: 7, AR267: 6, AR294: 6, AR226: 6, AR183: 5, AR269: 5, AR293: 5, AR258: 4, AR033: 4, AR268: 4, AR231: 4,		

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AR244:	4, AR283:	4,
AR243:	4, AR291:	3,
AR184:	3, AR290:	3,
AR299:	3, AR251:	3,
AR282:	3, AR310:	3,
AR296:	3, AR316:	2,
AR247:	2, AR089:	2,
AR052:	2, AR246:	2,
AR233:	2, AR186:	2,
AR237:	2, AR185:	2,
AR281:	2, AR313:	2,
AR096:	2, AR300:	1,
AR277:	1, AR060:	1,
AR179:	1, AR240:	1,
AR285:	1, AR055:	1,
AR061:	1, AR218:	1,
AR219:	1, AR280:	1,
AR259:	1, AR270:	1,
AR204:	1, AR312:	1,
AR202:	1, AR175:	0,
AR315:	0, AR177:	0,
AR275:	0, AR039:	0,
AR271:	0, AR192:	0,
AR206:	0, AR309:	0,
AR198:	0, AR248:	0,
AR205:	0, AR194:	0,
AR273:	0, AR213:	0,
AR249:	0, AR104:	0,
AR253:	0, AR265:	0

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114	HDPUN70	1096317	124	3 - 542	300	Ser-2 to Leu-9, Pro-26 to Tyr-36, His-40 to Asn-46, Ser-93 to Asn-102, Ser-107 to Leu-121, Ser-161 to Cys-169.	L0438: 1, H0547: 1, H0436: 1, L0731: 1, H0543: 1 and H0423: 1. AR313: 18, AR229: 13, AR241: 11, AR089: 11, AR192: 9, AR184: 9, AR271: 9, AR265: 8, AR299: 8, AR039: 8, AR096: 8, AR315: 8, AR238: 7, AR314: 7, AR263: 7, AR247: 7, AR226: 7, AR183: 7, AR182: 7, AR198: 7, AR312: 7, AR310: 6, AR248: 6, AR300: 6, AR202: 6, AR240: 6, AR269: 6, AR237: 6, AR281: 6, AR293: 6, AR244: 6, AR285: 6, AR185: 6, AR316: 5, AR270: 5, AR053: 5, AR296: 5, AR218: 5, AR258: 5, AR243: 5, AR206: 5, AR177: 5, AR286: 5, AR267: 5, AR231: 5, AR194: 5, AR292: 5, AR233: 4, AR249: 4, AR175: 4, AR280: 4, AR227: 4, AR295: 4, AR268: 4,		
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115	HHEPF57	1096422	125	831 - 2987	301	Lys-7 to Lys-13, Glu-135 to Leu-142, Asp-277 to Pro-282, Asn-284 to Phe-291, Glu-326 to Arg-332, Pro-411 to Val-418, Ala-457 to Pro-463, Thr-498 to Gly-503, Phe-528 to Gly-534, Tyr-610 to Lys-616,	AR312: 21, AR265: 20, AR309: 18, AR251: 18, AR273: 17, AR053: 15, AR052: 11, AR184: 11, AR219: 10, AR310: 10, AR243: 10, AR246: 9, AR104: 9, AR213: 9, AR218: 9, AR096: 8, AR198: 7, AR270: 7, AR186: 7, AR296: 6,				

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116	HFKFH60	1096718	126	402 - 2096	302	Asp-5 to Arg-11, Arg-47 to Val-53, Asp-62 to Ser-68, Asn-163 to Arg-170, Ala-217 to Glu-224,	AR205: 0, AR286: 0, AR283: 0, AR315: 0, AR244: 0, AR271: 0, AR280: 0 H0543: 18, L0438: 6, L0747: 5, L0439: 4, L0750: 4, L0779: 4, L0759: 4, L0592: 4, S0412: 4, H0156: 3, L0770: 3, L0758: 3, H0423: 3, H0402: 2, H0251: 2, L0809: 2, L0777: 2, H0542: 2, H0422: 2, H0624: 1, H0170: 1, S0114: 1, S0420: 1, S0007: 1, S6026: 1, H0351: 1, S6016: 1, H0013: 1, L0021: 1, H0575: 1, L0157: 1, T0010: 1, H0354: 1, S6028: 1, S0036: 1, H0038: 1, L0804: 1, L0787: 1, L0666: 1, H0658: 1 and L0742: 1.	8q22	216550, 259730
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Ala-258 to Gln-265, Asn-270 to Ser-276, Asn-286 to Thr-291, Glu-313 to Thr-332, Asp-337 to Cys-346, Arg-409 to Thr-414, Arg-450 to Val-461, Asn-475 to Thr-481, Thr-491 to Thr-506.	AR189: 12, AR191: 12, AR184: 9, AR267: 9, AR176: 9, AR178: 8, AR274: 8, AR052: 8, AR188: 7, AR180: 7, AR089: 7, AR247: 7, AR170: 6, AR215: 6, AR299: 6, AR181: 6, AR177: 6, AR235: 6, AR172: 5, AR273: 5, AR186: 5, AR271: 5, AR300: 5, AR310: 5, AR309: 5, AR236: 5, AR204: 5, AR060: 5, AR238: 5, AR201: 5, AR228: 5, AR196: 5, AR207: 4, AR316: 4, AR185: 4, AR289: 4, AR266: 4, AR312: 4, AR284: 4, AR288: 4, AR255: 4, AR225: 4, AR165: 4, AR161: 4, AR162: 4, AR171: 4, AR163: 4, AR168: 4, AR179: 4, AR166: 4, AR213: 4, AR313: 4, AR282: 4, AR055: 4, AR257: 4, AR164: 4, AR230: 4, AR253: 3, AR192: 3, AR294: 3
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AR033: 3, AR053: 3,									
AR263: 3, AR224: 3,									
AR245: 3, AR202: 3,									
AR239: 3, AR194: 3,									
AR277: 3, AR240: 3,									
AR206: 3, AR275: 3,									
AR233: 3, AR262: 3,									
AR251: 3, AR261: 3,									
AR221: 3, AR061: 3,									
AR291: 3, AR287: 3,									
AR169: 3, AR265: 3,									
AR285: 3, AR298: 3,									
AR311: 3, AR295: 3,									
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AR237: 3, AR198: 3,									
AR293: 3, AR226: 3,									
AR231: 3, AR217: 3,									
AR193: 3, AR203: 3,									
AR292: 2, AR234: 2,									
AR286: 2, AR223: 2,									
AR243: 2, AR296: 2,									
AR297: 2, AR096: 2,									
AR232: 2, AR212: 2,									
AR216: 2, AR219: 2,									
AR272: 2, AR039: 2,									
AR229: 2, AR195: 2,									
AR258: 2, AR218: 2,									
AR244: 2, AR199: 1,									
AR256: 1, AR211: 1,									
AR283: 1, AR227: 1,									

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		L0740: 2, L0749: 2, L0780: 2, L0601: 2, S0011: 2, H0668: 2, S0026: 2, H0543: 2, H0650: 1, L0785: 1, H0671: 1, H0661: 1, S0420: 1, S0356: 1, T0008: 1, H0580: 1, H0619: 1, H0645: 1, H0261: 1, H0549: 1, H0550: 1, S0222: 1, H0370: 1, H0486: 1, H0427: 1, H0036: 1, H0390: 1, S0474: 1, S0049: 1, H0546: 1, H0046: 1, H0457: 1, T0003: 1, S0050: 1, H0266: 1, H0271: 1, H0188: 1, H0687: 1, H0615: 1, H0428: 1, H0039: 1, L0483: 1, T0006: 1, H0644: 1, H0074: 1, H0674: 1, L0455: 1, H0163: 1, H0591: 1, H0264: 1, H0412: 1, H0059: 1, T0069: 1, T0042: 1, H0625: 1, S0150: 1, S0472: 1, H0649: 1, H0652: 1, S0344: 1,		
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117	HHBFK81	1096722	127	354 - 1811	303	His-1 to Gly-12, Arg-66 to Thr-71, Arg-107 to Val-118, Asn-132 to Thr-138	S0422: 1, S0426: 1, H0695: 1, L0598: 1, L0762: 1, L0640: 1, L0638: 1, L0796: 1, L0637: 1, L0772: 1, L0646: 1, L0800: 1, L0363: 1, L0767: 1, L0768: 1, L0650: 1, L0375: 1, L0784: 1, L0526: 1, L0384: 1, S0052: 1, S0053: 1, H0702: 1, S0374: 1, H0520: 1, H0547: 1, H0684: 1, H0660: 1, S0328: 1, S0330: 1, H0539: 1, S0380: 1, H0528: 1, S0013: 1, S0044: 1, H0555: 1, S0392: 1, H0626: 1, S0028: 1, L0744: 1, L0745: 1, H0595: 1, L0589: 1, L0590: 1, S0192: 1, S0242: 1, S0276: 1, S0196: 1, H0542: 1, H0422: 1 and H0506: 1.	AR225: 921, AR223: 766, AR296: 354, AR261: 331, AR287: 329, AR235: 324		
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				Glu-149 to Cys-167, Asp-191 to Glu-200, Phe-261 to Lys-268, Asp-317 to Val-324, Glu-374 to Lys-380, Glu-388 to Ala-398.	AR256: 318, AR297: 259, AR255: 237, AR289: 215, AR262: 209, AR196: 186, AR245: 168, AR236: 152, AR285: 141, AR288: 141, AR291: 131, AR266: 127, AR181: 113, AR258: 109, AR257: 107, AR294: 104, AR272: 96, AR286: 92, AR260: 92, AR211: 91, AR293: 81, AR180: 71, AR221: 70, AR210: 66, AR199: 65, AR222: 64, AR295: 64, AR219: 58, AR283: 57, AR224: 56, AR201: 54, AR178: 54, AR243: 54, AR200: 53, AR246: 53, AR165: 52, AR169: 52, AR188: 51, AR203: 51, AR218: 51, AR164: 50, AR166: 49, AR173: 48, AR170: 46, AR176: 45, AR191: 45, AR162: 44, AR316: 44, AR161: 43, AR163: 42, AR197: 42, AR189: 42, AR242: 41, AR193: 41, AR230: 40,	
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			AR207: 39, AR168: 39, AR039: 38, AR240: 37, AR183: 36, AR269: 35, AR171: 35, AR195: 33, AR290: 33, AR174: 33, AR274: 33, AR089: 32, AR238: 32, AR271: 31, AR270: 31, AR190: 31, AR055: 30, AR192: 29, AR267: 28, AR247: 27, AR175: 27, AR177: 27, AR179: 26, AR268: 26, AR231: 26, AR204: 24, AR311: 24, AR182: 22, AR212: 22, AR275: 21, AR313: 21, AR060: 21, AR214: 20, AR205: 20, AR215: 20, AR185: 19, AR239: 19, AR198: 18, AR228: 18, AR299: 16, AR300: 16, AR250: 16, AR234: 15, AR308: 15, AR282: 15, AR263: 12, AR264: 12, AR237: 11, AR253: 11, AR104: 11, AR277: 11, AR053: 10, AR213: 10, AR254: 10, AR172: 10, AR232: 9, AR061: 9, AR309: 9, AR096: 9, AR229: 9,						
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	AR184: 9, AR033: 9, AR226: 9, AR216: 8, AR217: 8, AR244: 8, AR312: 8, AR315: 8, AR292: 7, AR233: -7, AR052: 7, AR252: 6, AR284: 6, AR248: 5, AR273: 5, AR186: 5, AR298: 5, AR280: 5, AR314: 4, AR206: 4, AR249: 4, AR227: 4, AR265: 4, AR259: 3, AR310: 3, AR281: 1, AR202: 1, AR194: 1, AR251: 0, AR241: 0 L0747: 12, L0748: 9, L0740: 8, L0754: 8, L0749: 8, L0731: 8, L0438: 6, L0752: 6, L0809: 5, L0439: 5, L0756: 5, H0624: 4, H0620: 4, L0803: 4, L0783: 4, L0666: 4, L0663: 4, L0755: 4, S0354: 3, H0013: 3, H0123: 3, H0553: 3, S0426: 3, L0766: 3, L0774: 3, L0805: 3, S0152: 3, L0750: 3, L0779: 3, L0777: 3.	
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118	HARNY39	1097714	128	2 - 1225	304	Leu-64 to Glu-72, Asn-90 to Asp-98, Asn-154 to Val-176, Val-276 to Cys-282, Ser-323 to Gly-333, Pro-339 to Leu-353.	S0214: 1, H0615: 1, H0644: 1, H0674: 1, H0068: 1, H0163: 1, H0591: 1, H0551: 1, T0069: 1, S0438: 1, L0763: 1, L0772: 1, L0372: 1, L0764: 1, L0662: 1, L0767: 1, L0768: 1, L0650: 1, L0375: 1, L0784: 1, L0776: 1, L0655: 1, L0527: 1, L0384: 1, L0789: 1, L0790: 1, H0519: 1, H0435: 1, H0659: 1, H0658: 1, H0648: 1, S0328: 1, H0539: 1, H0522: 1, S0013: 1, H0696: 1, S0044: 1, S0406: 1, H0626: 1, S0037: 1, L0744: 1, L0751: 1, H0445: 1, L0589: 1, L0590: 1, S0242: 1, H0506: 1 and H0352: 1. AR314: 219, AR245: 218, AR315: 198, AR280: 195, AR281: 170, AR247: 68, AR292: 60, AR096: 54, AR283: 49, AR284: 44, AR298:	19p12	601843
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Gln-382 to Gly-393, Ser-397 to Arg-408.	39, AR285: 38, AR184: 34, AR259: 33, AR295: 33, AR300: 31, AR219: 30, AR256: 30, AR313: 29, AR291: 28, AR089: 28, AR266: 28, AR218: 27, AR231: 26, AR293: 25, AR240: 24, AR282: 24, AR033: 23, AR299: 23, AR183: 23, AR296: 23, AR286: 23, AR055: 23, AR268: 22, AR179: 22, AR289: 21, AR316: 21, AR294: 21, AR191: 20, AR198: 20, AR229: 20, AR188: 19, AR238: 19, AR232: 18, AR267: 18, AR175: 18, AR190: 17, AR177: 17, AR290: 17, AR165: 16, AR164: 16, AR277: 15, AR166: 15, AR237: 15, AR161: 15, AR162: 15, AR234: 15, AR258: 14, AR189: 14, AR163: 14, AR192: 14, AR269: 14, AR061: 14, AR185: 14, AR196: 13, AR182: 13, AR270: 13, AR226: 12, AR227: 11, AR215: 11, AR265:
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119	HIPDH20	1099283	129	3 - 869	305	His-14 to Gln-19, Gly-39 to Ala-47, Pro-58 to Thr-65, Tyr-168 to Lys-176, Gln-185 to Tyr-192, Glu-259 to Pro-265.	L0755: 1, L0596: 1, S0192: 1, S0194: 1, S0196: 1 and H0543: 1. AR281: 16, AR194: 15, AR206: 14, AR284: 14, AR315: 13, AR265: 13, AR310: 13, AR241: 13, AR244: 12, AR202: 12, AR205: 12, AR232: 12, AR061: 12, AR280: 12, AR309: 10, AR039: 10, AR289: 10, AR238: 10, AR052: 10, AR298: 10, AR282: 10, AR243: 10, AR096: 10, AR170: 9, AR246: 9, AR312: 9, AR192: 9, AR314: 9, AR215: 9, AR283: 9, AR268: 9, AR273: 9, AR089: 9, AR171: 9, AR299: 8, AR053: 8, AR213: 8, AR033: 8, AR277: 8, AR285: 8, AR291: 8, AR207: 8, AR183: 8, AR292: 8, AR168: 8, AR223: 8, AR184: 8, AR221: 8, AR060: 8, AR229: 8, AR182: 8, AR172: 8, AR316: 8, AR240: 8,	11p15.5	125852, 126452, 126452, 141900, 141900, 141900, 141900, 141900, 141900, 142000, 142000, 142200, 142250, 142270, 176730, 176730, 176730, 190020, 191290, 192500, 192500, 194071, 194071, 204500, 600856, 601680, 602631,
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	AR261: 5, AR216: 5, AR185: 5, AR219: 4, AR218: 4, AR195: 4, AR287: 4, AR199: 4, AR173: 4, AR242: 4, AR200: 4, AR236: 4, AR274: 4, AR257: 4, AR311: 4, AR297: 4, AR256: 4, AR191: 4, AR258: 4, AR189: 4, AR228: 4, AR179: 3, AR188: 3, AR201: 3, AR255: 3, AR190: 3, AR254: 3, AR262: 3, AR230: 3, AR174: 3, AR253: 3, AR210: 3, AR178: 3, AR203: 3, AR211: 2, AR260: 2, AR252: 1 H0305: 4, L0731: 3, L0581: 3, H0622: 2, H0059: 2, L0764: 2, L0766: 2, L0741: 2, L0740: 2, L0749: 2, H0423: 2, H0149: 1, H0159: 1, S0114: 1, H0255: 1, H0306: 1, H0402: 1, S0045: 1, H0351: 1, H0550: 1, H0441: 1, H0036: 1.
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Asp-197 to His-211, Ser-223 to Ala-236, Ser-247 to Asp-257.	AR286: 3, AR237: 2, AR186: 2, AR227: 2, AR183: 2, AR310: 2, AR206: 2, AR283: 2, AR052: 2, AR273: 2, AR226: 2, AR060: 2, AR204: 2, AR316: 2, AR061: 2, AR247: 2, AR089: 2, AR295: 2, AR267: 2, AR175: 1, AR241: 1, AR096: 1, AR313: 1, AR232: 1, AR055: 1, AR300: 1, AR177: 1, AR256: 1, AR240: 1, AR285: 1, AR244: 1, AR185: 1, AR293: 1, AR263: 1, AR033: 1, AR039: 1, AR294: 1, AR259: 1, AR292: 1, AR104: 1, AR218: 1, AR202: 0, AR281: 0, AR258: 0, AR253: 0, AR219: 0, AR315: 0, AR314: 0, AR198: 0, AR309: 0, AR271: 0, AR280: 0, AR248: 0, AR205: 0, AR243: 0, AR192: 0, AR179: 0	S0354: 2, H0059: 2.
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121	HWLED94	1099509	131	2 - 1624	307			L0803: 2, L0659: 2, L0809: 2, L0791: 2, L0748: 2, L0749: 2, S0358: 1, H0575: 1, H0085: 1, H0014: 1, H0051: 1, H0040: 1, L0773: 1, L0774: 1, L0375: 1, L0805: 1, L0527: 1, H0672: 1, S0380: 1, L0596: 1 and L0593: 1.		
								AR104:1359, AR233: 965, AR229: 896, AR186: 896, AR182: 797, AR298: 796, AR227: 794, AR234: 767, AR179: 751, AR237: 738, AR284: 712, AR061: 706, AR259: 668, AR238: 665, AR055: 664, AR314: 640, AR285: 610, AR185: 582, AR295: 564, AR060: 553, AR256: 538, AR300: 537, AR267: 519, AR258: 508, AR175: 503, AR292: 503, AR206: 501, AR226: 499, AR286:	2q37.3	

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[illegible]

122	HKAJ55	1100188	132	3 - 740	308	Pro-14 to Arg-25, Glu-29 to Cys-40, Gln-62 to Val-67, Glu-100 to Ser-107, Pro-117 to Ser-129, Ala-144 to Trp-157, Ser-173 to Gly-183.	H0264: 1, H0412: 1, H0100: 1, L0351: 1, H0561: 1, S0450: 1, H0538: 1, L0371: 1, L0775: 1, L0776: 1, L0659: 1, L0809: 1, L0663: 1, L0665: 1, S0053: 1, H0670: 1, H0672: 1, H0539: 1, H0518: 1, S0350: 1, L0786: 1, L0755: 1 and S0192: 1.		
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AR222: 2, AR291: 2,							
AR288: 2, AR216: 2,							
AR312: 2, AR238: 2,							
AR198: 2, AR224: 2,							
AR231: 2, AR186: 2,							
AR226: 2, AR266: 2,							
AR228: 1, AR240: 1,							
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AR290: 1, AR299: 1,							
AR061: 1, AR253: 1,							
AR239: 1, AR193: 1,							
AR089: 1, AR176: 1,							
AR292: 1, AR232: 1,							
AR178: 1, AR294: 1,							
AR316: 1, AR096: 1,							
AR274: 1, AR227: 1,							
AR217: 1, AR234: 1,							
AR271: 1, AR052: 1,							
AR247: 1, AR190: 1,							
AR060: 1, AR264: 1,							
AR313: 1, AR185: 1,							
AR213: 1, AR053: 1,							
AR283: 1, AR188: 1,							
AR221: 1, AR246: 1,							
AR175: 1, AR262: 1,							
AR196: 1, AR295: 1,							
AR260: 1, AR236: 1,							
AR233: 1, AR273: 1,							
AR191: 1, AR308: 1,							
AR258: 1, AR293: 1,							

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123	HCEPF75	1100345	133	1470 - 403	309	Arg-1 to Glu-6, Phe-17 to Gln-22, Asp-171 to Glu-177, Ser-267 to Cys-273, Pro-299 to Val-306, Arg-322 to Gly-345, Lys-348 to Arg-355.	AR244: 11, AR206: 8, AR186: 8, AR055: 4, AR204: 4, AR241: 4, AR273: 4, AR235: 4, AR271: 4, AR251: 4, AR033: 4, AR052: 4, AR310: 4, AR184: 4, AR243: 4, AR061: 3, AR183: 3, AR176: 3, AR291: 3, AR164: 3, AR165: 3, AR246: 3, AR166: 3, AR245: 3, AR298: 3, AR300: 3, AR299: 3, AR309: 3, AR248: 3, AR202: 3, AR162: 3, AR269: 3, AR053: 3, AR282: 3, AR182: 3, AR274: 2, AR259: 2, AR264: 2, AR180: 2, AR277: 2, AR228: 2, AR192: 2, AR236: 2, AR104: 2, AR247: 2, AR185: 2, AR286: 2, AR238: 2, AR289: 2, AR284: 2, AR312: 2, AR270: 2, AR257: 2, AR229: 2, AR205: 2, AR096: 2, AR292: 2, AR089: 2, AR201: 2, AR232: 2,	22q13.31- q13.33	250100, 250800, 250800
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AR267: 2, AR175: 2,							
AR161: 2, AR225: 2,							
AR283: 2, AR213: 2,							
AR168: 2, AR173: 2,							
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AR163: 2, AR181: 2,							
AR313: 2, AR290: 2,							
AR190: 2, AR261: 2,							
AR316: 2, AR198: 2,							
AR174: 2, AR293: 2,							
AR177: 2, AR268: 2,							
AR285: 2, AR227: 1,							
AR178: 1, AR231: 1,							
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AR223: 1, AR253: 1,							
AR230: 1, AR266: 1,							
AR294: 1, AR258: 1,							
AR287: 1, AR242: 1,							
AR203: 1, AR262: 1,							
AR193: 1, AR219: 1,							
AR217: 1, AR288: 1,							
AR275: 1, AR255: 1,							
AR239: 1, AR200: 1,							
AR194: 1, AR039: 1,							
AR297: 1, AR256: 1,							
AR234: 1, AR171: 1,							

124	HRDFA94	1100389	134	1 - 1065	310	Gly-1 to Asn-10, Val-18 to Lys-25, Leu-69 to Arg-78, Asn-83 to Pro-89, Thr-244 to Lys-258, Gly-293 to Gly-298.	AR308: 1, AR215: 1, AR172: 1, AR314: 1, AR263: 1, AR216: 1, AR169: 1, AR260: 1, AR218: 1, AR207: 1, AR281: 1, AR179: 1, AR197: 1, AR199: 0, AR254: 0, AR196: 0, AR214: 0, AR191: 0, AR224: 0, AR170: 0, AR252: 0, AR280: 0, AR188: 0, AR315: 0, AR210: 0, AR249: 0, AR212: 0, AR265: 0 H0052: 4, L0731: 2, H0624: 1, H0650: 1, H0351: 1, H0123: 1, T0023: 1, L0666: 1, H0144: 1, H0672: 1, H0539: 1, L0777: 1 and S0260: 1.	20p13	192340, 234200
							AR298: 285, AR055: 272, AR259: 267, AR227: 249, AR233: 234, AR226: 228, AR284: 218, AR294: 211, AR292: 206, AR286: 196, AR285: 194, AR234: 188, AR182: 180, AR231:		

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125	HSIFQ16	1100483	135	71 - 1483	311	Leu-35 to Lys-41, Leu-61 to Glu-68, Ser-153 to Gln-158, Asn-223 to Pro-228, Ala-259 to Phe-266, Pro-276 to Gly-283, Asp-292 to Phe-307, Ala-318 to Asp-336, Pro-348 to Leu-365, Ala-369 to Thr-393, Gln-398 to Gly-409, Ala-413 to Leu-419, Asn-441 to Ala-448.	AR176: 11, AR235: 11, AR181: 9, AR170: 8, AR225: 8, AR180: 7, AR173: 6, AR260: 6, AR224: 6, AR216: 6, AR222: 6, AR169: 6, AR214: 5, AR171: 5, AR215: 5, AR244: 5, AR217: 5, AR168: 5, AR223: 4, AR221: 4, AR172: 3, AR210: 1, AR211: 1 HO124: 1 and HO521: 1.	7q22	126650, 126650, 154276, 173360, 173360, 602136, 602136, 602136, 602447
						AR183: 8, AR312: 8, AR313: 8, AR194: 7, AR309: 7, AR273: 7, AR275: 6, AR248: 6, AR249: 6, AR274: 6, AR192: 6, AR213: 6, AR186: 6, AR263: 6, AR053: 6, AR246: 6, AR206: 6, AR265: 6, AR060: 5, AR204: 5, AR052: 5, AR310: 5, AR251: 5, AR253: 5, AR055: 5, AR277: 5, AR219: 5, AR282: 5, AR202: 4, AR185: 4, AR299: 4, AR229: 4, AR089: 4, AR243: 4,			

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126	HWLDY64	1100528	136	1178 - 390	312	Pro-118 to Tyr-123, Lys-187 to Val-192.	H0457: 2, H0539: 2, H0521: 2, L0744: 2, S0040: 1, H0583: 1, H0650: 1, H0484: 1, H0254: 1, S0356: 1, S0358: 1, H0592: 1, H0497: 1, H0257: 1, H0590: 1, H0318: 1, H0581: 1, H0024: 1, H0634: 1, T0041: 1, H0695: 1, H0529: 1, H0697: 1, S0378: 1, H0436: 1, L0743: 1, H0136: 1, H0543: 1, H0423: 1, H0422: 1 and H0677: 1. AR184: 8, AR248: 5, AR244: 4, AR270: 4, AR183: 4, AR268: 4, AR175: 3, AR282: 3, AR096: 3, AR182: 3, AR234: 3, AR310: 3, AR269: 2, AR238: 2, AR295: 2, AR265: 2, AR232: 2, AR284: 2, AR204: 2, AR229: 2, AR285: 2, AR274: 2, AR226: 1, AR237: 1, AR291: 1, AR060: 1, AR292: 1, AR281: 1,		
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127	HELB20	1100919	137	2707 - 2447	313	Arg-1 to Gly-6, Gln-23 to Ser-35, Ile-62 to Asp-72, His-74 to Ser-82.	L0774: 1, S0152: 1, L0758: 1 and L0608: 1. AR186: 8, AR194: 8, AR192: 7, AR241: 7, AR246: 6, AR055: 6, AR060: 6, AR263: 6, AR202: 6, AR052: 6, AR061: 6, AR269: 6, AR251: 6, AR206: 6, AR185: 5, AR265: 5, AR204: 5, AR053: 5, AR270: 5, AR243: 5, AR273: 5, AR267: 5, AR198: 5, AR268: 5, AR313: 5, AR253: 5, AR266: 4, AR247: 4, AR312: 4, AR310: 4, AR295: 4, AR213: 4, AR177: 4, AR282: 4, AR316: 4, AR286: 4, AR299: 4, AR033: 4, AR284: 4, AR298: 4, AR182: 4, AR183: 4, AR309: 4, AR249: 4, AR089: 4, AR289: 4, AR300: 4, AR205: 3, AR274: 3, AR229: 3, AR290: 3, AR096: 3, AR184: 3, AR285: 3, AR296: 3, AR275: 3,		
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128	HETJF84	1102857	138	2792 - 114	314	Pro-12 to Arg-27, Arg-29 to Phe-38, Tyr-42 to Thr-48, Val-52 to Ile-60, Gly-79 to Ser-89, Pro-91 to Asn-120, Ser-122 to Trp-127, Thr-129 to Gly-158, Pro-160 to Met-171, Asp-175 to Trp-181, Gly-192 to Asp-203, Pro-213 to Gln-220,	AR196: 6, AR202: 6, AR235: 6, AR275: 6, AR191: 5, AR174: 5, AR265: 5, AR184: 4, AR263: 4, AR189: 4, AR273: 4, AR186: 4, AR175: 4, AR161: 4, AR162: 4, AR163: 4, AR192: 4, AR246: 3, AR196: 3, AR188: 3, AR310: 3, AR268: 3, AR251: 3, AR176: 3,	1				

Pro-230 to Arg-237, Asp-260 to Asp-265, Asp-368 to Gln-375, Asp-407 to Arg-412, Glu-426 to Ile-432, Lys-446 to Ser-455, Pro-477 to Asn-484, Asp-502 to Asp-510, Arg-516 to Asn-527, Ser-540 to Asp-547, Asp-553 to Ser-560, Pro-562 to Cys-567, Pro-580 to Met-590, Asn-593 to Tyr-613, Gly-622 to Asn-628, Glu-636 to Tyr-646, Gln-648 to Tyr-657, Arg-663 to Gly-671, Glu-689 to Gly-695, Pro-703 to Asp-716, Pro-746 to Val-756.	AR269: 3, AR221: 3, AR311: 3, AR270: 3, AR272: 3, AR288: 3, AR225: 3, AR290: 3, AR177: 3, AR254: 3, AR287: 3, AR232: 3, AR206: 2, AR170: 2, AR282: 2, AR284: 2, AR181: 2, AR296: 2, AR245: 2, AR240: 2, AR291: 2, AR312: 2, AR298: 2, AR299: 2, AR294: 2, AR055: 2, AR183: 2, AR262: 2, AR061: 2, AR257: 2, AR060: 2, AR199: 2, AR096: 2, AR295: 2, AR178: 2, AR217: 2, AR224: 2, AR214: 2, AR171: 2, AR204: 2, AR247: 2, AR203: 2, AR201: 2, AR089: 2, AR168: 2, AR316: 2, AR266: 2, AR165: 2, AR172: 2, AR207: 2, AR233: 2, AR285: 2, AR039: 2, AR300: 2, AR286: 2, AR238: 2, AR185: 2, AR033: 2, AR166: 2, AR309: 2,
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AR205:	1, AR274:	1,							
AR258:	1, AR236:	1,							
AR182:	1, AR219:	1,							
AR289:	1, AR226:	1,							
AR218:	1, AR281:	1,							
AR200:	1, AR292:	1,							
AR222:	1, AR229:	1,							
AR271:	1, AR231:	1,							
AR255:	1, AR252:	1,							
AR256:	1, AR241:	1,							
AR242:	1, AR223:	1,							
AR261:	1, AR211:	1,							
AR104:	1, AR180:	1,							
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AR259:	0, AR260:	0,							
AR264:	0, AR244:	0,							
AR193:	0, AR249:	0,							
AR216:	0, AR194:	0,							
AR195:	0, AR250:	0,							
AR237:	0, AR213:	0,							
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AR212:	0, AR308:	0,							
AR280:	0, AR315:	0,							

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129	HE2DL46	1102938	139	34 - 1458	315	His-8 to Gly-18, Gly-73 to Asn-79, Leu-81 to Trp-86, Ser-91 to Gln-99, Asn-101 to Phe-118, Cys-190 to Phe-199, Cys-222 to Ser-241, Ser-262 to Phe-280, Arg-298 to Tyr-303, Asp-318 to Thr-324, Ser-330 to Asp-336, Ile-339 to Trp-344, Phe-348 to Gln-357, Glu-359 to Gly-368, Asp-381 to Val-395, Thr-438 to Asp-446.	AR249: 4, AR166: 3, AR250: 3, AR282: 3, AR198: 3, AR253: 3, AR216: 3, AR212: 2, AR180: 2, AR309: 2, AR183: 2, AR215: 1, AR217: 1, AR168: 1, AR165: 1, AR161: 1, AR312: 1, AR299: 1, AR272: 1, AR240: 1, AR172: 1, AR089: 1, AR291: 1, AR277: 1, AR308: 1, AR211: 1, AR181: 1, AR096: 1, AR266: 1, AR189: 1, AR246: 1, AR287: 1, AR194: 1, AR281: 1, AR313: 1, AR255: 1,				

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130	HDPSJ02	1103186	140	2 - 766	316	Asp-10 to Ala-15, Leu-42 to Phe-52, His-110 to Leu-115, Gln-141 to Arg-150, Phe-168 to Arg-174, Pro-231 to Tyr-237.	H0570: 1, H0051: 1, H0598: 1, S0036: 1, H0144: 1, L0759: 1, S0031: 1, L0592: 1 and S0106: 1.	20p13	192340, 234200
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131	HE6FT04	1103911	141	147 - 1460	317	Pro-49 to Thr-66, Arg-80 to Gly-88, Ala-92 to Glu-105, Lys-141 to Asp-147, Pro-180 to Gly-185, Glu-188 to Gly-207, Arg-238 to Asn-246, Glu-266 to Ala-276, Gly-303 to Gly-318, Pro-333 to Gly-345, Ser-424 to Arg-429.	H0617: 1, S0366: 1, S0142: 1, L0768: 1, S0374: 1, H0690: 1, H0672: 1, H0522: 1, H0555: 1, L0743: 1, L0749: 1, L0777: 1 and L0731: 1. AR184: 4, AR244: 4, AR282: 3, AR241: 3, AR182: 3, AR277: 2, AR243: 2, AR251: 2, AR292: 2, AR284: 2, AR273: 2, AR290: 2, AR183: 2, AR269: 2, AR270: 2, AR313: 2, AR229: 2, AR293: 1, AR312: 1, AR267: 1, AR039: 1, AR247: 1, AR175: 1, AR274: 1, AR296: 1, AR289: 1, AR213: 1, AR310: 1, AR240: 1, AR299: 1, AR286: 1, AR089: 1, AR238: 1, AR249: 1, AR096: 1, AR316: 1, AR300: 1, AR283: 1, AR234: 1, AR237: 1, AR298: 1, AR246: 1, AR291: 0, AR266: 0, AR052: 0, AR061: 0.		
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132	HETAQ37	1103950	142	1 - 1380	318		AR227: 0, AR271: 0, AR314: 0, AR259: 0, AR055: 0, AR309: 0, AR294: 0, AR205: 0, AR060: 0, AR256: 0, AR198: 0, AR053: 0, AR275: 0, AR219: 0, AR281: 0, AR186: 0, AR258: 0, AR192: 0, AR263: 0, AR202: 0, AR177: 0, AR280: 0, AR265: 0, AR033: 0, AR231: 0, AR204: 0, AR226: 0, AR206: 0, AR218: 0, AR248: 0, AR268: 0, AR233: 0, AR185: 0, AR295: 0, AR179: 0, AR104: 0, AR315: 0 H0052: 1, L0163: 1, H0644: 1, H0100: 1, S0150: 1, L0517: 1, H0539: 1, H0521: 1, H0555: 1, L0748: 1, L0750: 1, L0588: 1 and L0594: 1.	12q14.1- 12q15	181430, 600698, 600698, 600698,
							AR202:1350, AR206:1040, AR204:1013, AR244: 782, AR246: 572,		

600698, 600808, 602116		AR194: 539, AR205: 535, AR233: 512, AR232: 485, AR241: 476, AR177: 466, AR281: 463, AR230: 437, AR300: 426, AR039: 423, AR185: 423, AR055: 413, AR227: 407, AR186: 403, AR198: 399, AR033: 396, AR271: 393, AR060: 390, AR228: 377, AR283: 375, AR061: 365, AR226: 361, AR179: 353, AR259: 340, AR104: 314, AR299: 314, AR273: 313, AR294: 312, AR315: 311, AR239: 307, AR180: 295, AR178: 294, AR192: 293, AR310: 289, AR280: 282, AR295: 281, AR275: 281, AR237: 278, AR174: 277, AR243: 275, AR234: 269, AR247: 261, AR274: 260, AR254: 257, AR282: 252
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AR297: 145, AR309: 144, AR164: 142, AR235: 142, AR096: 142, AR212: 141, AR267: 140, AR286: 135, AR277: 135, AR195: 133, AR183: 127, AR219: 121, AR291: 112, AR261: 112, AR190: 111, AR313: 110, AR272: 108, AR289: 101, AR169: 101, AR214: 95, AR311: 93, AR223: 92, AR266: 91, AR211: 91, AR176: 90, AR296: 83, AR203: 81, AR260: 80, AR191: 80, AR298: 77, AR170: 75, AR222: 73, AR269: 73, AR308: 69, AR264: 69, AR189: 67, AR224: 65, AR173: 64, AR199: 64, AR184: 62, AR284: 60, AR171: 58, AR270: 58, AR168: 58, AR210: 57, AR172: 57, AR268: 57, AR225: 55, AR216: 54, AR290: 54, AR217: 52, AR196: 51, AR188: 49, AR215: 47,									
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133	HCENW09	1104186	143	2 - 673	319	Trp-16 to Leu-21, Asp-89 to Asn-96.	AR200: 45, AR221: 42 H0046: 44, H0135: 10, H0539: 10, L0455: 7, L0750: 5, S0010: 3, L0456: 3, L0779: 3, L0005: 2, L0663: 2, S0328: 2, L0746: 2, L0747: 2, L0777: 2, H0624: 1, S0116: 1, H0208: 1, L0717: 1, H0549: 1, H0333: 1, H0013: 1, L0021: 1, L0157: 1, T0006: 1, H0652: 1, L0515: 1, L0666: 1, H0144: 1, H0696: 1, L0439: 1 and L0745: 1.	15	
							AR263: 6, AR265: 4, AR310: 4, AR269: 4, AR312: 3, AR313: 3, AR296: 3, AR182: 3, AR244: 3, AR183: 3, AR266: 3, AR253: 3, AR270: 3, AR089: 3, AR249: 2, AR247: 2, AR299: 2, AR096: 2, AR314: 2, AR033: 2, AR292: 2, AR295: 2, AR277: 2, AR285: 2, AR282: 2, AR284: 2,		

AR293:	2, AR175:	2,
AR290:	2, AR184:	2,
AR238:	2, AR298:	2,
AR248:	2, AR061:	2,
AR229:	2, AR289:	2,
AR177:	2, AR267:	2,
AR316:	2, AR268:	2,
AR286:	1, AR192:	1,
AR309:	1, AR227:	1,
AR294:	1, AR060:	1,
AR233:	1, AR053:	1,
AR232:	1, AR237:	1,
AR315:	1, AR055:	1,
AR226:	1, AR231:	1,
AR213:	1, AR206:	1,
AR271:	1, AR243:	1,
AR291:	1, AR234:	1,
AR258:	1, AR104:	1,
AR300:	1, AR259:	1,
AR186:	1, AR185:	1,
AR275:	1, AR219:	1,
AR205:	1, AR256:	1,
AR281:	1, AR039:	1,
AR283:	1, AR240:	1,
AR218:	0, AR251:	0,
AR280:	0, AR241:	0,
AR204:	0, AR179:	0,
AR194:	0, AR273:	0,
AR246:	0, AR052:	0,
AR202:	0, AR198:	0,

134	HAMFW05	1104784	144	504 - 2246	320	Arg-2 to Pro-9, Glu-14 to Arg-22, Pro-24 to Pro-31, Gln-41 to Gly-46, Thr-51 to Leu-56, Asp-88 to Trp-93, Thr-108 to Arg-114, Pro-149 to Glu-154, Ser-181 to Glu-196, Gln-234 to Pro-240.	AR274: 0 H0052: 4, H0014: 2, H0549: 1, H0194: 1, L0163: 1, T0010: 1, H0083: 1, H0622: 1, H0135: 1, L0438: 1, H0520: 1, H0547: 1, H0682: 1, S0028: 1, L0439: 1, L0745: 1 and L0777: 1. AR313: 21, AR249: 15, AR296: 15, AR229: 12, AR183: 12, AR238: 12, AR089: 10, AR292: 10, AR293: 10, AR258: 10, AR269: 10, AR182: 9, AR234: 9, AR096: 9, AR263: 9, AR298: 9, AR270: 9, AR226: 9, AR291: 8, AR248: 8, AR184: 8, AR219: 8, AR299: 8, AR265: 8, AR218: 8, AR312: 8, AR285: 7, AR267: 7, AR247: 7, AR240: 7, AR294: 7, AR233: 7, AR268: 6, AR300: 6, AR231: 6, AR316: 6, AR033: 6, AR310: 6, AR290: 6, AR241: 6,		
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									AR237: 6, AR104: 6, AR175: 6, AR289: 6, AR259: 6, AR280: 6, AR177: 5, AR314: 5, AR286: 5, AR295: 5, AR256: 5, AR185: 5, AR227: 5, AR192: 5, AR282: 5, AR198: 5, AR186: 5, AR253: 4, AR060: 4, AR315: 4, AR179: 4, AR284: 4, AR052: 4, AR053: 4, AR213: 4, AR232: 4, AR309: 3, AR277: 3, AR266: 3, AR271: 3, AR204: 3, AR251: 3, AR281: 3, AR039: 3, AR274: 3, AR275: 3, AR283: 2, AR273: 2, AR061: 2, AR243: 2, AR055: 2, AR205: 1, AR206: 1, AR202: 1, AR194: 1, AR244: 1, AR246: 0 H0521: 6, L0757: 5, L0749: 4, L0779: 4, H0620: 3, S0040: 2, S0342: 2, H0305: 2, S0418: 2, S0132: 2, H0551: 2, H0412: 2.
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	Glu-133 to Gly-139, Ser-173 to Phe-186, Thr-194 to Ser-205, Ala-233 to Gly-238, Ser-281 to Ser-294, Pro-302 to Thr-313.	AR266: 4, AR269: 4, AR313: 4, AR273: 4, AR312: 4, AR292: 3, AR283: 3, AR213: 3, AR289: 3, AR309: 3, AR314: 3, AR243: 3, AR286: 3, AR248: 3, AR270: 3, AR052: 3, AR284: 3, AR039: 2, AR310: 2, AR053: 2, AR182: 2, AR282: 2, AR265: 2, AR202: 2, AR294: 2, AR096: 2, AR175: 2, AR186: 2, AR240: 2, AR231: 2, AR033: 2, AR295: 2, AR238: 2, AR249: 2, AR206: 2, AR316: 2, AR234: 2, AR244: 2, AR290: 2, AR275: 2, AR300: 2, AR299: 2, AR277: 2, AR253: 2, AR104: 2, AR315: 2, AR267: 2, AR298: 2, AR237: 2, AR229: 2, AR293: 2, AR089: 2, AR185: 2, AR268: 2, AR205: 1, AR285: 1, AR247: 1, AR246: 1, AR256: 1, AR177: 1,
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AR179: 1, AR280: 1, AR227: 1, AR226: 1, AR192: 1, AR258: 1, AR218: 1, AR259: 1, AR055: 1, AR219: 1, AR060: 1, AR281: 1, AR061: 1, AR271: 1, AR194: 1, AR251: 1, AR198: 0, AR233: 0, AR204: 0, AR232: 0 L0748: 9, L0754: 6, L0659: 5, H0046: 4, H0494: 4, H0556: 3, S0046: 3, H0644: 3, L0764: 3, H0521: 3, L0750: 3, L0759: 3, L0595: 3, L0603: 3, H0295: 2, H0254: 2, S0045: 2, H0370: 2, H0024: 2, H0264: 2, H0623: 2, S0142: 2, S0344: 2, S0002: 2, L0768: 2, L0783: 2, L0664: 2, H0520: 2, L0740: 2, L0596: 2, H0294: 1, T0049: 1, H0255: 1, H0662: 1, H0638: 1, S0356: 1, S0358: 1, H0580: 1, S0132: 1, H0619: 1,									
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136	HDTEJ03	1104936	146	686 - 2260	322	Ser-40 to Glu-51,	H0393: 1, S6022: 1, H0587: 1, H0331: 1, H0013: 1, H0575: 1, H0590: 1, S0474: 1, H0123: 1, H0050: 1, S0051: 1, S0368: 1, H0031: 1, H0617: 1, H0598: 1, H0634: 1, H0063: 1, H0268: 1, H0059: 1, H0100: 1, H0625: 1, H0396: 1, H0509: 1, H0641: 1, H0647: 1, H0646: 1, H0649: 1, H0652: 1, L0763: 1, L0646: 1, L0662: 1, L0363: 1, L0649: 1, L0389: 1, L0774: 1, L0653: 1, L0655: 1, L0658: 1, L0636: 1, L0517: 1, L0790: 1, L0666: 1, S0374: 1, S0310: 1, H0547: 1, S0328: 1, H0518: 1, S0152: 1, S0013: 1, S0044: 1, H0436: 1, L0743: 1, L0749: 1, L0752: 1, L0581: 1, L0593: 1 and S0456: 1.	AR309: 9, AR263: 8,
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Arg-84 to Pro-96, Gly-119 to His-127, Val-132 to Cys-141, Pro-194 to Asn-203, Lys-231 to Tyr-239, Gln-245 to Asn-252, Val-272 to Tyr-280, Pro-287 to Cys-297, Gly-324 to Trp-334, Pro-338 to Cys-343, Pro-346 to Trp-361, Cys-384 to Gln-395, Arg-397 to Ser-406, Pro-409 to Asp-420, Lys-428 to Asp-441, Ala-511 to His-516.	AR310: AR265: AR238: AR184: AR292: AR313: AR295: AR253: AR291: AR226: AR186: AR285: AR284: AR290: AR089: AR233: AR316: AR294: AR293: AR247: AR267: AR248: AR033: AR232: AR104: AR275: AR185: AR246: AR229: AR234:	7, AR312: 6, AR266: 5, AR202: 5, AR270: 4, AR299: 4, AR269: 4, AR096: 4, AR052: 4, AR286: 3, AR183: 3, AR282: 3, AR182: 3, AR268: 3, AR271: 3, AR213: 3, AR053: 3, AR251: 3, AR060: 3, AR289: 3, AR296: 3, AR298: 3, AR274: 2, AR273: 2, AR240: 2, AR175: 2, AR277: 2, AR231: 2, AR237: 2, AR300: 1, AR177:	6, 5, 5, 5, 4, 4, 4, 4, 3, 3, 3, 3, 3, 3, 3, 3, 3, 3, 3, 3, 3, 2, 2, 2, 2, 2, 2, 2, 2, 1, 1,
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AR039: 1, AR259: 1, AR258: 1, AR061: 1, AR055: 1, AR249: 1, AR283: 1, AR256: 1, AR314: 1, AR243: 1, AR315: 1, AR244: 1, AR194: 1, AR198: 1, AR219: 1, AR179: 1, AR192: 1, AR218: 0, AR227: 0, AR280: 0, AR205: 0, AR206: 0, AR241: 0 H0549: 7, L0665: 6, L0751: 6, L0439: 5, H0620: 3, L0803: 3, L0777: 3, L0601: 3, H0483: 2, H0486: 2, H0309: 2, H0687: 2, L0774: 2, L0657: 2, L0659: 2, L0809: 2, L0666: 2, L0438: 2, H0520: 2, H0593: 2, H0658: 2, L0602: 2, H0555: 2, H0624: 1, H0686: 1, H0295: 1, H0656: 1, S0282: 1, H0255: 1, S0354: 1, H0580: 1, H0619: 1, H0618: 1, H0581: 1, S0049: 1, H0052: 1,									
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137	HTTSA02	1105584	147	706 - 2	323				H0562: 1, H0012: 1, H0083: 1, S0250: 1, H0428: 1, L0483: 1, H0135: 1, S0038: 1, H0494: 1, L0640: 1, L0638: 1, L0637: 1, L0771: 1, L0662: 1, L0805: 1, L0655: 1, L0629: 1, L0368: 1, L0789: 1, L0663: 1, H0519: 1, H0682: 1, H0670: 1, H0521: 1, H0522: 1, L0740: 1, L0779: 1, H0667: 1 and H0543: 1.		107776, 138079, 138079, 139191, 180104, 203740, 601472, 601649
									AR273: 15, AR274: 12, AR309: 11, AR313: 10, AR243: 10, AR251: 9, AR310: 9, AR053: 9, AR312: 8, AR271: 8, AR263: 7, AR213: 7, AR265: 7, AR052: 7, AR282: 7, AR219: 6, AR218: 6, AR247: 6, AR175: 6, AR096: 6, AR248: 5, AR183: 5, AR240: 5, AR089: 5, AR249: 5, AR299: 5, AR275: 5, AR266: 5, AR241: 5, AR316: 4	7p14	

AR292:	4, AR280:	4,
AR184:	4, AR194:	4,
AR060:	4, AR177:	4,
AR269:	4, AR186:	4,
AR185:	4, AR253:	4,
AR256:	4, AR295:	4,
AR204:	4, AR206:	4,
AR198:	3, AR104:	3,
AR039:	3, AR205:	3,
AR314:	3, AR290:	3,
AR268:	3, AR291:	3,
AR293:	3, AR246:	3,
AR270:	3, AR277:	3,
AR294:	3, AR283:	3,
AR258:	3, AR285:	3,
AR289:	3, AR315:	3,
AR179:	2, AR231:	2,
AR238:	2, AR055:	2,
AR232:	2, AR300:	2,
AR298:	2, AR286:	2,
AR033:	2, AR267:	2,
AR234:	2, AR061:	2,
AR296:	2, AR182:	2,
AR281:	2, AR227:	2,
AR284:	2, AR226:	2,
AR237:	2, AR259:	2,
AR233:	1, AR192:	1,
AR202:	1, AR244:	1,
AR229:	0	
S0264:	28, S0254:	27,

138	H7TSA19	1105585	148	509 - 3	324		S0258: 9, S0256: 1, S0252: 1 and L0534: 1.	AR274: 18, AR251: 15, AR273: 14, AR313: 13, AR053: 12, AR309: 12, AR310: 11, AR312: 11, AR271: 10, AR247: 10, AR243: 9, AR263: 9, AR096: 9, AR219: 9, AR290: 9, AR218: 8, AR282: 8, AR265: 8, AR213: 8, AR266: 8, AR089: 8, AR183: 7, AR315: 7, AR052: 7, AR280: 7, AR314: 7, AR175: 7, AR240: 6, AR060: 6, AR316: 6, AR299: 6, AR291: 6, AR289: 6, AR268: 6, AR292: 6, AR286: 6, AR248: 6, AR184: 6, AR267: 6, AR285: 5, AR104: 5, AR256: 5, AR293: 5, AR182: 5, AR295: 5, AR269: 5, AR253: 5, AR198: 5, AR296: 5, AR185: 5, AR275: 5, AR283: 5, AR244: 5, AR294: 5, AR249: 5, AR270: 5,	7p14	107776, 138079, 138079, 139191, 180104, 203740, 601472, 601649
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139	HMWCU63	1105747	149	46 - 375	325	Asn-50 to Pro-55, Ser-78 to Asp-95.	AR284: 5, AR039: 5, AR186: 5, AR298: 4, AR177: 4, AR205: 4, AR277: 4, AR192: 4, AR179: 4, AR231: 3, AR281: 3, AR300: 3, AR226: 3, AR033: 3, AR229: 3, AR055: 3, AR232: 3, AR234: 3, AR061: 3, AR238: 3, AR259: 3, AR237: 3, AR246: 3, AR258: 3, AR204: 2, AR233: 2, AR202: 2, AR206: 2, AR194: 1, AR227: 1, AR241: 1 S0254: 84, S0264: 44, S0258: 30, S0256: 9, S0252: 1 and L0534: 1.					103950, 120580, 131440, 139130, 142680, 176260, 190450, 200990, 216950, 600228, 600414,
							AR314: 55, AR280: 53, AR219: 52, AR315: 43, AR218: 43, AR291: 40, AR270: 36, AR052: 36, AR266: 31, AR268: 30, AR290: 29, AR183: 27, AR269: 26, AR298: 25, AR289: 25, AR313: 23, AR296: 22, AR267: 22, AR053: 22, AR281: 22, AR293: 21, AR292: 21,	12p13				

AR294: 19, AR285: 19, AR175: 18, AR312: 17, AR309: 17, AR182: 16, AR184: 16, AR256: 15, AR259: 15, AR284: 14, AR231: 14, AR258: 14, AR299: 13, AR213: 13, AR263: 13, AR310: 13, AR179: 12, AR316: 12, AR096: 11, AR295: 11, AR238: 11, AR060: 11, AR237: 11, AR265: 11, AR286: 10, AR234: 9, AR177: 9, AR089: 8, AR300: 7, AR229: 7, AR249: 7, AR240: 6, AR282: 6, AR248: 5, AR232: 5, AR247: 5, AR033: 5, AR226: 5, AR241: 5, AR253: 4, AR055: 4, AR185: 4, AR192: 3, AR061: 3, AR251: 3, AR243: 3, AR233: 3, AR277: 3, AR227: 2, AR283: 2, AR246: 2, AR186: 1, AR205: 1, AR204: 0, AR244: 0, AR202: 0, AR274: 0, AR198: 0, AR206: 0, AR273: 0,	600618, 602096
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140	HWADF26	1105857	150	300 - 947	326	Arg-9 to Pro-15, Ala-63 to Asp-71, Asn-75 to Asp-86, Glu-102 to Cys-110, Asn-115 to Ser-123.	AR039: 0, AR271: 0, AR275: 0 H0656: 2, H0341: 2, H0038: 2, H0616: 2, L0766: 2, H0518: 2, L0748: 2, H0542: 2, H0543: 2, H0221: 1, H0158: 1, H0036: 1, H0571: 1, H0109: 1, H0560: 1, H0561: 1, L0763: 1, L0662: 1, L0794: 1, L0775: 1, L0655: 1, L0663: 1, H0648: 1, H0525: 1 and H0423: 1.		
							AR264: 35, AR197: 22, AR170: 19, AR168: 16, AR311: 15, AR212: 15, AR169: 15, AR171: 14, AR265: 14, AR207: 14, AR223: 14, AR195: 14, AR225: 13, AR308: 13, AR214: 12, AR217: 12, AR193: 11, AR216: 11, AR196: 11, AR165: 11, AR161: 11, AR162: 11, AR210: 11, AR163: 11, AR261: 11, AR245: 11, AR164: 10, AR235: 10, AR250: 10, AR180: 10,		

[illegible]

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AR219:	4, AR182:	4,							
AR294:	4, AR247:	4,							
AR267:	4, AR234:	4,							
AR033:	4, AR179:	4,							
AR089:	4, AR277:	3,							
AR230:	3, AR310:	3,							
AR039:	3, AR316:	3,							
AR296:	3, AR258:	3,							
AR238:	3, AR300:	3,							
AR244:	3, AR293:	3,							
AR228:	3, AR299:	3,							
AR060:	3, AR256:	3,							
AR052:	3, AR229:	3,							
AR285:	3, AR104:	3,							
AR286:	3, AR055:	3,							
AR233:	3, AR275:	2,							
AR218:	2, AR185:	2,							
AR237:	2, AR226:	2,							
AR231:	2, AR283:	2,							
AR204:	2, AR251:	2,							
AR273:	2, AR186:	2,							
AR061:	2, AR232:	2,							
AR206:	1, AR248:	1,							
AR249:	1, AR227:	1,							
AR284:	1, AR194:	1,							
AR298:	1, AR241:	1,							
AR259:	1, AR202:	0,							
AR292:	0, AR281:	0,							
AR315:	0, AR314:	0,							

141	HDABT82	1106030	151	470 - 1288	327	Gln-48 to Lys-64, Glu-175 to Thr-183.	AR280: 0 H0581: 2, S0114: 1 and L0362: 1. AR249: 40, AR268: 31, AR290: 26, AR270: 19, AR184: 17, AR096: 16, AR267: 16, AR183: 15, AR194: 14, AR269: 14, AR206: 13, AR263: 12, AR229: 12, AR205: 12, AR281: 11, AR246: 11, AR299: 11, AR202: 10, AR192: 10, AR271: 9, AR175: 9, AR039: 9, AR182: 9, AR243: 9, AR253: 8, AR251: 8, AR310: 8, AR177: 8, AR198: 7, AR248: 7, AR295: 7, AR316: 7, AR265: 7, AR282: 7, AR315: 7, AR283: 7, AR234: 7, AR291: 7, AR241: 7, AR247: 7, AR186: 7, AR292: 6, AR273: 6, AR280: 6, AR204: 6, AR238: 6, AR313: 6, AR240: 6, AR266: 6, AR300: 6, AR289: 6, AR274: 6, AR277: 6, AR226: 5,		
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142	HDPMQ42	1106192	152	214 - 963	328	Asn-1 to Ser-9, Lys-26 to Ser-31, Ala-41 to Pro-48.	AR248: 46, AR104: 38, AR186: 33, AR251: 31, AR055: 28, AR061: 26, AR266: 25, AR033: 24, AR185: 24, AR249: 23, AR243: 23, AR296: 21, AR198: 21, AR206: 20,	12p13	103950, 120580, 131440, 139130, 142680, 176260, 190450,
							H0090: 2, L0771: 2, L0773: 2, L0776: 2, L0655: 2, L0809: 2, H0555: 2, L0756: 2, L0362: 2, H0624: 1, S0358: 1, S0360: 1, H0637: 1, H0645: 1, H0392: 1, H0497: 1, H0486: 1, S0280: 1, H0042: 1, H0318: 1, H0014: 1, H0375: 1, H0644: 1, H0068: 1, S0142: 1, L0769: 1, L0646: 1, L0662: 1, L0651: 1, L0792: 1, L0664: 1, S0374: 1, H0658: 1, S0380: 1, L0754: 1, L0779: 1, L0780: 1, L0753: 1, L0758: 1, H0445: 1, H0595: 1, L0597: 1, H0653: 1, S0424: 1 and H0506: 1.		

AR204: 20, AR241: 20, AR283: 18, AR233: 18, AR289: 18, AR298: 18, AR271: 17, AR089: 17, AR192: 16, AR253: 16, AR194: 16, AR273: 15, AR234: 15, AR316: 15, AR227: 15, AR247: 15, AR229: 14, AR060: 14, AR281: 14, AR282: 14, AR291: 14, AR313: 14, AR240: 13, AR202: 13, AR237: 13, AR300: 13, AR290: 13, AR292: 13, AR286: 13, AR184: 12, AR314: 12, AR315: 12, AR205: 12, AR285: 12, AR096: 12, AR039: 12, AR280: 12, AR284: 12, AR246: 11, AR293: 11, AR182: 11, AR232: 11, AR274: 11, AR263: 10, AR294: 10, AR183: 10, AR226: 10, AR275: 10, AR295: 9, AR231: 9, AR269: 9, AR277: 9, AR310: 9, AR270: 8, AR256: 8, AR244: 8, AR299: 8, AR268: 8, AR238: 8, AR267: 8,	200990, 216950, 600228, 600414, 600618, 602096
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143	HHFHQ39	1106209	153	1 - 2124	329	Gly-1 to Asp-6, Ser-16 to Thr-24, Asn-47 to Gly-54.	AR265: 7, AR175: 7, AR259: 7, AR258: 7, AR177: 7, AR309: 7, AR213: 6, AR053: 6, AR179: 6, AR312: 6, AR052: 5, AR218: 4, AR219: 3 H0521: 10, H0522: 4, H0580: 2, S0214: 2, L0750: 2, H0486: 1, L0021: 1, H0251: 1, H0309: 1, H0263: 1, H0271: 1, S0003: 1, H0068: 1, H0090: 1, S0144: 1, L0803: 1, L0804: 1, L0749: 1, L0777: 1 and L0759: 1.	10q25.3- q26.1	246600, 263700, 600035, 601969, 601969
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AR277: 1, AR033: 1,									
AR316: 1, AR089: 1,									
AR300: 1, AR175: 1,									
AR234: 1, AR184: 1,									
AR229: 1, AR232: 1,									
AR246: 1, AR194: 1,									
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AR198: 1, AR226: 1,									
AR206: 1, AR289: 1,									
AR244: 1, AR268: 1,									
AR233: 1, AR284: 0,									
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144	HAFAQ92	1106214	154	90 - 2258	330			H0622: 32, H0039: 15, L0646: 7, H0506: 5, S0330: 4, S0358: 3, L0803: 2, L0806: 2, L0783: 2, S0328: 2, L0599: 2, L0362: 2, H0624: 1, S0376: 1, H0489: 1, H0036: 1, H0050: 1, H0040: 1, H0623: 1, H0386: 1, L0761: 1, L0772: 1, L0373: 1, L0372: 1, L0642: 1, L0771: 1, L0649: 1, L0776: 1, L0659: 1, L0747: 1, L0777: 1, S0456: 1 and S0446: 1.		
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145	HLHCR16	1106227	155	1 - 2901	331				AR183: 21, AR177: 16, AR269: 15, AR270: 12, AR175: 11, AR192: 10, AR182: 9 AR268: 9				

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146	HTGDN55	1106269	156	116 - 217	332	Leu-1 to Asp-6, Ser-13 to Thr-22.	AR264: 0, AR199: 0, AR287: 0, AR203: 0, AR297: 0, AR161: 0, AR280: 0, AR315: 0, AR168: 0, AR239: 0, AR162: 0, AR262: 0 L0754: 14, L0777: 13, H0553: 10, L0600: 7, L0748: 6, L0803: 4, L0749: 4, UNKWN: 4, H0624: 3, S0280: 3, S0126: 3, L0747: 3, S0282: 2, H0024: 2, H0030: 2, H0031: 2, H0040: 2, L0438: 2, S0028: 2, L0743: 2, L0596: 2, L0603: 2, S0212: 1, H0270: 1, H0244: 1, H0427: 1, H0251: 1, H0309: 1, S0338: 1, S0340: 1, S0250: 1, H0252: 1, H0039: 1, L0143: 1, H0038: 1, L0659: 1, L0565: 1, H0593: 1, H0684: 1, H0518: 1, S0390: 1, S0260: 1 and H0506: 1.		
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147	H7MCB42	H7MCB42R P00A	157	2 - 103	333					H0555: 1, S0392: 1, S0322: 1, L0748: 1, L0740: 1, L0745: 1, L0749: 1, L0756: 1, L0757: 1 and S0242: 1. AR226: 8, AR227: 6, AR232: 6, AR263: 5, AR238: 4, AR237: 4, AR275: 3, AR061: 3, AR282: 3, AR271: 3, AR233: 3, AR184: 3, AR251: 2, AR205: 2, AR246: 2, AR198: 2, AR231: 2, AR299: 2, AR267: 2, AR292: 2, AR298: 2, AR060: 2, AR285: 1, AR240: 1, AR186: 1, AR310: 1, AR234: 1, AR286: 1, AR253: 1, AR039: 1, AR277: 1, AR316: 1, AR312: 1, AR053: 1, AR270: 1, AR192: 1, AR268: 1, AR096: 1, AR089: 1, AR293: 1, AR241: 1, AR296: 1, AR185: 1, AR281: 1, AR052: 1, AR289: 1, AR244: 1, AR179: 1, AR177: 1 AR202: 1
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148	H7TWD35	H7TWD35F	158	1 - 408	334	AR259: 1, AR290: 1, AR104: 1, AR313: 1, AR033: 1, AR300: 0, AR229: 0, AR291: 0, AR280: 0, AR256: 0, AR274: 0, AR314: 0, AR258: 0, AR265: 0, AR219: 0, AR315: 0, AR055: 0, AR213: 0, AR273: 0, AR283: 0, AR175: 0, AR294: 0, AR194: 0, AR284: 0, AR295: 0, AR266: 0, AR249: 0, AR206: 0, AR247: 0, AR309: 0, AR218: 0, AR248: 0		
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AR192:	0, AR296:	0,							
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AR268:	0, AR039:	0,							
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149	HBCMB46	HBCMB46 R	159	3 - 203	335		AR177: 20, AR280: 15, AR314: 14, AR315: 13, AR269: 12, AR183: 11, AR270: 10, AR271: 9, AR175: 8, AR192: 7, AR274: 7, AR243: 7, AR204: 7, AR299: 7, AR268: 6, AR184: 6, AR266: 6, AR194: 5, AR281: 5, AR273: 5, AR179: 5, AR182: 5, AR247: 5, AR267: 5, AR293: 5, AR275: 5, AR263: 4, AR198: 4, AR206: 4, AR292: 4, AR291: 4, AR313: 4, AR290: 4, AR096: 4, AR316: 4, AR240: 4, AR205: 4, AR295: 4, AR300: 3, AR089: 3, AR186: 3, AR289: 3, AR294: 3, AR277: 3, AR248: 3, AR238: 3, AR282: 3, AR060: 3, AR265: 3, AR256: 3, AR185: 3, AR296: 3, AR052: 3, AR246: 2, AR298: 2, AR234: 2, AR251: 2, AR053: 2,		
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150	HCEBJ34	HCEBJ34R	160	76 - 252	336			AR219: 2, AR231: 2, AR285: 2, AR232: 2, AR259: 2, AR237: 2, AR213: 2, AR312: 2, AR229: 2, AR286: 2, AR227: 2, AR033: 2, AR253: 1, AR218: 1, AR061: 1, AR244: 1, AR284: 1, AR055: 1, AR233: 1, AR226: 1, AR258: 1, AR283: 1, AR104: 1, AR310: 1, AR309: 0, AR249: 0, AR202: 0, AR039: 0, AR241: 0		
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151	HCUBO07	HCUBO07R	161	1 - 141	337				AR241: 1, AR185: 1, AR247: 1, AR291: 1, AR248: 1, AR234: 1, AR283: 1, AR275: 0, AR270: 0, AR271: 0, AR218: 0, AR061: 0, AR253: 0, AR184: 0, AR052: 0, AR259: 0, AR192: 0, AR213: 0, AR267: 0, AR243: 0, AR300: 0, AR177: 0, AR182: 0, AR244: 0, AR231: 0, AR229: 0, AR281: 0, AR313: 0, AR273: 0, AR258: 0, AR240: 0, AR314: 0, AR295: 0, AR055: 0, AR285: 0, AR290: 0, AR233: 0, AR227: 0, AR060: 0, AR237: 0, AR299: 0, AR226: 0, AR179: 0, AR219: 0, AR256: 0, AR246: 0, AR198: 0, AR286: 0		
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152	HDPWO71	HDPWO71 R	162	2 - 241	338		AR266: 0, AR198: 0, AR315: 0, AR289: 0, AR039: 0, AR249: 0, AR204: 0, AR231: 0, AR205: 0, AR275: 0, AR206: 0, AR194: 0, AR253: 0, AR243: 0, AR213: 0, AR251: 0, AR273: 0		
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153	HDPYN86	HDPYN86R	163	1 - 150	339				11, AR300: 11, AR179: 11, AR282: 10, AR204: 10, AR234: 10, AR269: 10, AR183: 9, AR185: 8, AR089: 8, AR267: 7, AR060: 6, AR182: 5, AR186: 5, AR231: 5, AR033: 4, AR293: 4, AR295: 4, AR061: 4, AR284: 4, AR226: 3, AR229: 3, AR285: 3, AR294: 3, AR232: 3, AR291: 3, AR104: 2, AR298: 2, AR277: 2, AR238: 2, AR283: 2, AR237: 2, AR286: 2, AR233: 2, AR292: 2, AR296: 2, AR258: 2, AR289: 1, AR259: 1, AR256: 1, AR227: 1, AR266: 1, AR314: 1, AR218: 0, AR280: 0, AR219: 0, AR315: 0, AR281: 0		
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154	HDQED69	HDQED69R	164	2 - 127	340			AR263: 11, AR253: 10, AR265: 9, AR052: 7, AR310: 7, AR309: 7, AR249: 7, AR312: 7, AR275: 6, AR243: 6, AR248: 6, AR246: 6, AR251: 6, AR192: 6, AR053: 6, AR096: 6, AR213: 5, AR183: 5, AR313: 5, AR271: 5, AR273: 5, AR240: 4, AR247: 4, AR039: 4, AR184: 3, AR299: 3, AR206: 3, AR198: 3, AR316: 3, AR055: 3, AR315: 3, AR282: 3, AR186: 3, AR231: 2, AR185: 2, AR177: 2, AR277: 2, AR089: 2, AR294: 2, AR270: 2, AR218: 2, AR234: 2, AR280: 2, AR182: 2, AR219: 2, AR292: 2,			

155	HDQHW94	HDQHW94 R	165	1 - 138	341		AR295: 2, AR298: 2, AR237: 2, AR060: 2, AR285: 2, AR194: 2, AR179: 1, AR175: 1, AR061: 1, AR033: 1, AR205: 1, AR286: 1, AR281: 1, AR314: 1, AR238: 1, AR229: 1, AR104: 1, AR300: 1, AR241: 1, AR267: 1, AR293: 1, AR258: 1, AR226: 1, AR266: 1, AR289: 1, AR283: 1, AR256: 1, AR233: 1, AR232: 1, AR244: 0, AR274: 0, AR269: 0, AR290: 0, AR259: 0, AR284: 0, AR202: 0, AR296: 0, AR227: 0, AR291: 0, AR204: 0		
							AR096: 160, AR249: 135, AR253: 135, AR183: 118, AR268: 115, AR179: 98, AR270: 92, AR177: 85, AR175: 83, AR184: 82, AR248: 74, AR240: 74, AR263: 69, AR269: 68, AR290: 67, AR267: 61, AR247:		

						58, AR265: 50, AR218: 46, AR234: 42, AR251: 36, AR219: 35, AR055: 33, AR231: 33, AR229: 31, AR316: 31, AR299: 31, AR238: 31, AR300: 29, AR313: 29, AR182: 28, AR293: 26, AR310: 24, AR282: 24, AR284: 23, AR089: 23, AR237: 21, AR294: 20, AR291: 20, AR259: 19, AR258: 18, AR226: 18, AR298: 17, AR285: 17, AR295: 16, AR292: 16, AR232: 15, AR033: 14, AR185: 14, AR296: 14, AR289: 13, AR256: 12, AR061: 12, AR060: 11, AR233: 11, AR213: 10, AR286: 9, AR281: 9, AR283: 9, AR315: 9, AR280: 8, AR053: 8, AR314: 8, AR266: 7, AR227: 7, AR052: 5, AR277: 5, AR312: 4, AR194: 4, AR204: 3, AR309: 3, AR186: 3, AR274: 3, AR271: 2, AR206: 1, AR104: 1, AR244:
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156	HDQPO42	HDQPO42R	166	159 - 329	342			1, AR202: 1, AR246: 1, AR241: 1, AR198: 0, AR192: 0, AR275: 0, AR039: 0, AR273: 0, AR243: 0, AR205: 0		
								AR253: 84, AR249: 68, AR213: 62, AR248: 58, AR263: 56, AR265: 47, AR270: 47, AR268: 44, AR184: 43, AR269: 40, AR052: 37, AR313: 36, AR290: 36, AR312: 35, AR096: 32, AR182: 31, AR053: 30, AR310: 29, AR309: 29, AR251: 26, AR267: 23, AR291: 22, AR285: 18, AR284: 18, AR247: 16, AR298: 15, AR316: 15, AR296: 15, AR089: 14, AR299: 14, AR183: 13, AR271: 11, AR234: 11, AR246: 11, AR244: 11, AR194: 11, AR266: 10, AR238: 10, AR226: 9, AR286: 9, AR231: 9, AR202: 8, AR243: 8, AR289: 8, AR177: 8, AR039: 8, AR186: 8, AR229: 8,		

157	HEMEP27	HEMEP27R	167	3 - 401	343	Pro-7 to Lys-13.	AR282: 7, AR237: 7, AR055: 7, AR060: 7, AR205: 7, AR275: 7, AR175: 6, AR185: 6, AR300: 6, AR204: 6, AR240: 6, AR192: 6, AR315: 6, AR314: 6, AR280: 6, AR198: 5, AR206: 5, AR241: 5, AR033: 5, AR232: 5, AR292: 5, AR273: 5, AR295: 4, AR061: 4, AR277: 4, AR218: 3, AR219: 3, AR293: 3, AR274: 3, AR294: 3, AR281: 3, AR227: 3, AR179: 3, AR233: 3, AR104: 2, AR259: 2, AR283: 2, AR258: 1, AR256: 1		
							AR291: 62, AR266: 60, AR256: 47, AR219: 46, AR259: 45, AR295: 38, AR218: 38, AR292: 38, AR284: 36, AR296: 35, AR285: 34, AR289: 34, AR298: 34, AR286: 32, AR293: 31, AR104: 28, AR258: 27, AR183: 27.		

	AR269: 27, AR294: 26, AR175: 24, AR270: 23, AR182: 21, AR309: 21, AR268: 20, AR186: 20, AR290: 19, AR179: 18, AR184: 18, AR312: 18, AR275: 17, AR248: 17, AR213: 17, AR273: 17, AR310: 16, AR265: 16, AR313: 16, AR033: 16, AR246: 16, AR316: 15, AR263: 15, AR271: 15, AR253: 14, AR053: 14, AR240: 14, AR244: 14, AR052: 13, AR205: 13, AR231: 13, AR251: 13, AR274: 13, AR267: 13, AR249: 13, AR177: 12, AR237: 12, AR185: 11, AR282: 11, AR238: 11, AR243: 11, AR060: 11, AR283: 10, AR247: 10, AR229: 10, AR241: 9, AR192: 9, AR096: 9, AR198: 9, AR234: 9, AR089: 9, AR206: 8, AR194: 8, AR039: 8, AR299: 8, AR204: 7, AR226: 7, AR300: 6, AR202: 6, AR232: 5.
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158	HKGBC43	HKGBC43R	168	1 - 273	344	Phe-2 to Trp-7, Thr-29 to Glu-39, Leu-48 to Gln-54.	AR053: 20, AR309: 19, AR271: 19, AR265: 19, AR263: 14, AR310: 13, AR244: 10, AR312: 8, AR243: 6, AR206: 5, AR273: 5, AR282: 5, AR052: 5, AR213: 4, AR275: 3, AR060: 3, AR205: 3, AR283: 3, AR314: 3, AR293: 3, AR089: 3, AR316: 2, AR291: 2, AR315: 2, AR227: 2, AR253: 2, AR232: 2, AR280: 2, AR185: 1, AR175: 1, AR177: 1, AR270: 1, AR061: 1, AR289: 1, AR096: 1, AR033: 1, AR240: 1, AR286: 1, AR259: 1, AR281: 1, AR299: 1, AR241: 1, AR294: 1, AR247: 1, AR313: 1, AR186: 1, AR274: 1, AR204: 1,	AR061: 5, AR055: 5, AR277: 4, AR227: 3, AR233: 3, AR281: 1, AR314: 0, AR315: 0, AR280: 0		
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159	HLICD96	HLICD96R	169	2 - 277	345		AR246: 1, AR300: 1, AR256: 1, AR184: 1, AR277: 0, AR249: 0, AR251: 0, AR183: 0, AR284: 0, AR104: 0, AR238: 0, AR055: 0, AR269: 0, AR295: 0, AR194: 0, AR182: 0, AR218: 0, AR237: 0, AR039: 0, AR298: 0, AR258: 0, AR233: 0, AR290: 0, AR229: 0, AR202: 0, AR226: 0, AR219: 0, AR267: 0, AR296: 0, AR179: 0, AR234: 0, AR285: 0, AR192: 0, AR292: 0		
							AR313: 61, AR241: 58, AR281: 52, AR192: 51, AR194: 50, AR198: 49, AR315: 45, AR039: 42, AR089: 41, AR096: 41, AR202: 41, AR246: 40, AR229: 40, AR234: 40, AR244: 40, AR206: 39, AR280: 39, AR204: 38, AR243: 38, AR314: 37, AR205: 36, AR238: 36, AR265: 36, AR300: 35,		

[illegible]

							AR298: 13, AR275: 12, AR290: 11			
160	HLJBC28	HLJBC28R P00E	170	339 - 713	346	Asp-1 to Ser-7, Gln-16 to Arg-21, Leu-34 to His-54, Glu-62 to Asp-71.	AR313: 20, AR184: 11, AR269: 10, AR293: 10, AR229: 10, AR182: 10, AR258: 10, AR219: 9, AR089: 9, AR218: 9, AR270: 9, AR183: 9, AR299: 9, AR296: 8, AR238: 8, AR226: 8, AR314: 7, AR268: 7, AR096: 7, AR292: 7, AR247: 7, AR259: 6, AR315: 6, AR285: 6, AR241: 6, AR053: 6, AR290: 6, AR267: 6, AR175: 5, AR294: 5, AR312: 5, AR240: 5, AR280: 5, AR234: 5, AR300: 5, AR316: 5, AR192: 5, AR052: 5, AR286: 5, AR282: 5, AR289: 5, AR060: 5, AR298: 5, AR227: 5, AR233: 5, AR291: 5, AR237: 4, AR177: 4, AR185: 4, AR231: 4, AR266: 4, AR277: 4, AR033: 4, AR295: 4,			

161	HL YEB70	HL YEB70R	171	1 - 153	347	AR284: 4, AR265: 4, AR244: 4, AR256: 3, AR263: 3, AR309: 3, AR249: 3, AR213: 3, AR232: 3, AR179: 3, AR310: 3, AR039: 3, AR194: 3, AR251: 3, AR274: 2, AR243: 2, AR186: 2, AR273: 2, AR283: 2, AR253: 2, AR205: 2, AR055: 1, AR061: 1, AR281: 1, AR104: 1, AR248: 1, AR275: 1, AR202: 1, AR246: 1, AR206: 1, AR198: 0, AR271: 0, AR204: 0 H0580: 1, H0575: 1, H0375: 1, H0271: 1, S0142: 1 and L0748: 1. AR096: 203, AR213: 161, AR249: 159, AR253: 139, AR240: 130, AR177: 124, AR268: 109, AR179: 106, AR183: 96, AR290: 89, AR053: 87, AR184: 86, AR270: 84, AR175: 84, AR247: 82, AR263: 80, AR248: 75, AR267:		
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162	HMCBK34	HMCBK34 R	172	2 - 136	348	AR213: 824, AR246: 752, AR096: 375, AR244: 358, AR240: 355, AR249: 349, AR053: 329, AR039: 273, AR253: 269, AR198: 259, AR192: 249, AR271: 233, AR290: 227, AR202: 215, AR052: 210, AR243: 210, AR241: 208, AR194: 203, AR268: 200, AR179: 150, AR183: 145, AR177: 140, AR270: 138, AR273: 125, AR275: 124, AR263: 122, AR184: 117, AR247: 116, AR274: 107, AR313: 104, AR269: 103, AR316: 102, AR175: 101, AR248: 100, AR205: 99, AR206: 97, AR267: 93	5, AR186: 5, AR227: 3, AR104: 2, AR219: 1, AR218: 1, AR281: 1, AR315: 1, AR280: 0, AR314: 0			
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163	HMKDT37	HMKDT37 R	173	1 - 144	349				AR055: 93, AR265: 80, AR251: 79, AR300: 74, AR204: 71, AR310: 70, AR312: 66, AR234: 66, AR309: 64, AR282: 60, AR299: 58, AR185: 57, AR060: 44, AR089: 43, AR061: 43, AR182: 42, AR186: 40, AR293: 36, AR284: 35, AR231: 32, AR104: 27, AR285: 25, AR229: 24, AR033: 23, AR298: 23, AR294: 23, AR291: 22, AR226: 21, AR258: 20, AR237: 20, AR283: 19, AR295: 19, AR259: 19, AR296: 17, AR238: 16, AR232: 15, AR286: 14, AR289: 14, AR292: 14, AR266: 14, AR233: 11, AR277: 11, AR256: 8, AR227: 5, AR281: 2, AR219: 2, AR218: 2, AR315: 2, AR314: 1, AR280: 1		
									AR265: 3, AR194: 3, AR204: 2, AR269: 2, AR266: 2, AR282: 2, AR229: 2, AR291: 2,		

AR096: 1, AR295: 1,					
AR244: 1, AR263: 1,					
AR267: 1, AR247: 1,					
AR270: 1, AR298: 1,					
AR033: 1, AR039: 1,					
AR234: 1, AR312: 1,					
AR177: 1, AR213: 1,					
AR286: 1, AR089: 1,					
AR310: 1, AR240: 1,					
AR251: 1, AR233: 1,					
AR205: 1, AR277: 1,					
AR274: 1, AR285: 1,					
AR185: 1, AR316: 1,					
AR290: 1, AR060: 1,					
AR226: 1, AR061: 0,					
AR246: 0, AR249: 0,					
AR241: 0, AR281: 0,					
AR198: 0, AR186: 0,					
AR300: 0, AR294: 0,					
AR253: 0, AR299: 0,					
AR184: 0, AR258: 0,					
AR313: 0, AR248: 0,					
AR256: 0, AR238: 0,					
AR231: 0, AR275: 0,					
AR289: 0, AR055: 0,					
AR309: 0, AR183: 0,					
AR284: 0, AR052: 0,					
AR175: 0, AR179: 0,					
AR202: 0, AR283: 0,					
AR292: 0, AR237: 0,					

164	HMPTG96	HMPTG96R P00E	174	1 - 561	350			AR227: 0, AR273: 0, AR259: 0, AR314: 0, AR271: 0, AR206: 0, AR232: 0, AR218: 0, AR315: 0, AR243: 0, AR053: 0		
								AR039: 39, AR194: 37, AR246: 35, AR202: 34, AR281: 34, AR205: 34, AR315: 31, AR244: 29, AR310: 28, AR192: 28, AR206: 28, AR263: 28, AR053: 28, AR198: 27, AR052: 25, AR243: 25, AR265: 25, AR280: 24, AR241: 24, AR309: 24, AR104: 24, AR273: 23, AR204: 23, AR213: 22, AR312: 22, AR033: 21, AR089: 21, AR314: 20, AR271: 20, AR282: 19, AR096: 18, AR275: 18, AR274: 18, AR295: 18, AR300: 18, AR285: 18, AR186: 17, AR240: 17, AR299: 17, AR316: 17, AR283: 16, AR313: 16, AR060: 15, AR284: 15, AR251: 15, AR247: 14,		

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H0656: 2, H0341: 2, S0358: 2, S0007: 2, H0575: 2, L0471: 2, H0266: 2, S0003: 2, H0615: 2, H0616: 2, L0598: 2, L0763: 2, L0769: 2, L0521: 2, L0803: 2, L0775: 2, L0655: 2, L0532: 2, H0144: 2, S0126: 2, H0682: 2, H0518: 2, L0740: 2, L0749: 2, L0757: 2, L0758: 2, H0595: 2, L0361: 2, S0026: 2, H0542: 2, H0171: 1, H0556: 1, H0221: 1, H0158: 1, H0222: 1, S0040: 1, T0049: 1, S0134: 1, H0657: 1, H0638: 1, H0125: 1, L0005: 1, S0045: 1, S0278: 1, H0438: 1, H0574: 1, H0036: 1, S0010: 1, H0318: 1, H0421: 1, L0738: 1, H0544: 1, L0157: 1, H0571: 1, S0388: 1, H0109: 1, H0628: 1, L0055: 1, H0090: 1, H0634: 1.					
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165	HMTBX02	HMTBX02	RP00	175	24 - 131	351	Tyr-7 to His-13, Ser-18 to Trp-25, Thr-31 to Gln-36.	H0412: 1, H0102: 1, H0494: 1, H0560: 1, H0561: 1, S0150: 1, H0130: 1, S0422: 1, H0529: 1, L0764: 1, L0794: 1, L0649: 1, L0776: 1, L0657: 1, L0545: 1, L0529: 1, L0541: 1, L0788: 1, L0791: 1, L0663: 1, L0665: 1, H0684: 1, H0648: 1, H0672: 1, H0525: 1, S0013: 1, L0756: 1, L0786: 1, L0755: 1, H0445: 1, S0308: 1, L0608: 1, S0192: 1, S0194: 1, H0423: 1 and S0424: 1, AR263: 14, AR265: 10, AR053: 8, AR052: 8, AR310: 8, AR309: 7, AR312: 6, AR313: 6, AR194: 5, AR213: 5, AR251: 5, AR282: 3, AR269: 3, AR270: 3, AR192: 3, AR183: 3, AR243: 3, AR249: 3, AR198: 3, AR184: 3, AR039: 3, AR271: 3, AR268: 3, AR096: 2,		
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								AR202: 2, AR229: 2, AR316: 2, AR205: 2, AR277: 2, AR299: 2, AR300: 2, AR283: 2, AR253: 2, AR285: 2, AR295: 2, AR267: 2, AR177: 2, AR186: 2, AR231: 2, AR293: 2, AR089: 1, AR033: 1, AR290: 1, AR240: 1, AR219: 1, AR234: 1, AR226: 1, AR238: 1, AR241: 1, AR104: 1, AR182: 1, AR060: 1, AR185: 1, AR247: 1, AR289: 1, AR227: 1, AR284: 1, AR218: 1, AR061: 1, AR233: 1, AR244: 1, AR179: 1, AR248: 1, AR175: 1, AR280: 1, AR206: 1, AR246: 0, AR275: 0, AR294: 0, AR256: 0, AR296: 0, AR258: 0, AR273: 0, AR259: 0, AR274: 0, AR281: 0, AR237: 0, AR232: 0, AR055: 0, AR292: 0, AR291: 0, AR314: 0, AR315: 0, AR298: 0,
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166	HMXAB55	HMXAB55 R	176	1 - 138	352	Asn-10 to Asp-17.	AR266: 0, AR204: 0 H0656: 2, H0341: 2, H0038: 2, H0616: 2, L0766: 2, H0518: 2, L0748: 2, H0542: 2, H0543: 2, H0221: 1, H0158: 1, H0036: 1, H0571: 1, H0109: 1, H0560: 1, H0561: 1, L0763: 1, L0662: 1, L0794: 1, L0775: 1, L0655: 1, L0663: 1, H0648: 1, H0525: 1 and H0423: 1.		
							AR273: 11, AR251: 10, AR309: 8, AR274: 7, AR313: 7, AR310: 7, AR312: 6, AR263: 6, AR266: 6, AR271: 6, AR052: 5, AR184: 5, AR053: 4, AR290: 4, AR282: 4, AR265: 4, AR291: 4, AR213: 4, AR253: 3, AR267: 3, AR247: 3, AR206: 3, AR268: 3, AR243: 3, AR285: 3, AR096: 3, AR060: 3, AR289: 3, AR292: 3, AR316: 3, AR089: 3, AR286: 2,		

167	HNFO84	HNFO84R	177	3 - 419	353			AR240: 2, AR284: 2, AR296: 2, AR269: 2, AR183: 2, AR185: 2, AR294: 2, AR270: 2, AR033: 2, AR182: 2, AR175: 2, AR299: 2, AR275: 2, AR249: 2, AR295: 2, AR298: 2, AR293: 2, AR186: 2, AR283: 2, AR277: 2, AR218: 2, AR104: 2, AR234: 1, AR300: 1, AR226: 1, AR061: 1, AR179: 1, AR232: 1, AR256: 1, AR229: 1, AR177: 1, AR055: 1, AR219: 1, AR238: 1, AR314: 1, AR280: 1, AR246: 1, AR194: 1, AR258: 1, AR233: 1, AR248: 1, AR039: 1, AR259: 0, AR315: 0, AR237: 0, AR244: 0, AR202: 0, AR204: 0, AR231: 0, AR205: 0, AR198: 0, AR227: 0, AR192: 0		AR248: 574, AR253: 374, AR251: 287,	
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168	HNTRS86	HNTRS86R	178	1 - 270	354	Leu-2 to Arg-8, Ser-65 to Glu-71.	AR313: 40, AR241: 29, AR096: 24, AR312: 23, AR258: 22, AR218: 21, AR247: 21, AR089: 21, AR183: 20, AR292: 19, AR299: 19, AR229: 18, AR052: 17, AR219: 17, AR265: 16, AR192: 16, AR263: 16, AR238: 15, AR300: 14, AR259: 14, AR314: 14, AR240: 14, AR033: 14, AR309: 14, AR177: 14, AR293: 14, AR315: 13, AR310: 13,	50, AR060: 48, AR237: 47, AR226: 42, AR258: 40, AR286: 38, AR053: 38, AR213: 38, AR300: 37, AR284: 36, AR204: 34, AR089: 34, AR299: 32, AR283: 31, AR232: 31, AR309: 31, AR282: 29, AR052: 29, AR233: 24, AR310: 24, AR265: 24, AR312: 24, AR055: 23, AR277: 22, AR227: 21, AR263: 21, AR314: 7, AR281: 6, AR315: 6, AR280: 5		
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AR053: 12, AR316: 12,									
AR198: 12, AR213: 12,									
AR256: 11, AR295: 11,									
AR280: 11, AR185: 11,									
AR231: 11, AR294: 10,									
AR175: 10, AR233: 10,									
AR226: 10, AR253: 10,									
AR249: 10, AR274: 10,									
AR271: 9, AR234: 9,									
AR275: 8, AR179: 8,									
AR186: 8, AR248: 8,									
AR237: 8, AR243: 8,									
AR104: 7, AR281: 7,									
AR282: 7, AR060: 7,									
AR296: 7, AR277: 7,									
AR204: 7, AR283: 7,									
AR285: 7, AR273: 7,									
AR298: 6, AR267: 6,									
AR291: 6, AR227: 6,									
AR270: 5, AR251: 5,									
AR290: 5, AR184: 5,									
AR039: 5, AR206: 4,									
AR266: 4, AR286: 4,									
AR269: 4, AR232: 4,									
AR182: 4, AR268: 3,									
AR284: 3, AR205: 3,									
AR289: 3, AR246: 3,									
AR061: 2, AR055: 2,									
AR244: 1, AR194: 1,									
AR202: 1									

169	HOEAC11	HOEAC11R	179	94 - 267	355	AR309: 31, AR052: 29, AR265: 27, AR312: 23, AR053: 21, AR270: 18, AR310: 18, AR313: 18, AR213: 18, AR263: 17, AR315: 16, AR314: 15, AR274: 15, AR269: 14, AR273: 14, AR243: 13, AR253: 12, AR280: 12, AR184: 12, AR296: 11, AR271: 10, AR290: 10, AR292: 10, AR291: 10, AR192: 10, AR182: 10, AR281: 9, AR186: 9, AR206: 9, AR268: 9, AR285: 9, AR284: 8, AR294: 8, AR293: 8, AR316: 8, AR060: 8, AR246: 8, AR202: 8, AR286: 7, AR205: 7, AR198: 7, AR248: 7, AR194: 7, AR249: 7, AR275: 7, AR267: 7, AR298: 7, AR244: 7, AR096: 6, AR039: 6, AR183: 6, AR104: 6, AR295: 6, AR175: 5, AR299: 5, AR266: 5, AR089: 5, AR289: 5,		
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170	HOFNH58	HOFNH58R	180	1 - 438	356		AR258: 5, AR251: 5, AR218: 5, AR033: 5, AR185: 5, AR219: 5, AR238: 5, AR055: 4, AR259: 4, AR240: 4, AR241: 4, AR282: 4, AR300: 4, AR179: 4, AR229: 4, AR204: 4, AR277: 3, AR177: 3, AR061: 3, AR247: 3, AR231: 3, AR237: 3, AR234: 3, AR232: 2, AR283: 2, AR233: 2, AR256: 2, AR227: 2, AR226: 2		
							AR270: 3, AR266: 3, AR184: 2, AR274: 2, AR282: 2, AR269: 2, AR285: 2, AR192: 2, AR310: 2, AR294: 2, AR286: 2, AR289: 2, AR229: 2, AR277: 2, AR312: 2, AR268: 2, AR253: 2, AR296: 2, AR298: 1, AR292: 1, AR231: 1, AR291: 1, AR198: 1, AR183: 1, AR053: 1, AR248: 1, AR241: 1, AR182: 1,		

[illegible]

171	HOHAL20	HOHAL20R	181	74 - 331	357	Pro-32 to Tyr-53.	AR263: 3, AR241: 2, AR194: 2, AR265: 2, AR184: 2, AR266: 2, AR244: 2, AR243: 2, AR295: 2, AR282: 1, AR096: 1, AR267: 1, AR247: 1, AR277: 1, AR290: 1, AR312: 1, AR232: 1, AR298: 1, AR310: 1, AR275: 1, AR289: 1, AR233: 1, AR238: 1, AR175: 1, AR291: 1, AR182: 1, AR313: 1, AR293: 1, AR270: 1, AR269: 1, AR089: 1, AR231: 1, AR033: 1, AR268: 1, AR294: 1, AR284: 1, AR316: 1, AR060: 1, AR061: 1, AR186: 1, AR227: 0, AR299: 0, AR285: 0, AR055: 0, AR251: 0, AR248: 0, AR234: 0, AR226: 0, AR296: 0, AR273: 0, AR177: 0, AR281: 0, AR202: 0, AR258: 0, AR259: 0, AR286: 0, AR253: 0, AR205: 0,		
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172	HS2AO61	HS2AO61R	182	21 - 290	358	Gln-43 to Asp-53.	AR229: 0, AR300: 0, AR240: 0, AR219: 0, AR218: 0, AR039: 0, AR206: 0, AR271: 0, AR204: 0, AR246: 0, AR237: 0, AR314: 0, AR185: 0, AR198: 0, AR183: 0, AR192: 0, AR256: 0, AR283: 0, AR053: 0, AR309: 0, AR179: 0, AR315: 0, AR104: 0, AR292: 0, AR213: 0, AR052: 0, AR249: 0, AR274: 0, AR280: 0		
							AR285: 6, AR269: 6, AR243: 6, AR270: 6, AR263: 6, AR265: 6, AR202: 5, AR266: 5, AR271: 5, AR313: 5, AR251: 5, AR182: 5, AR194: 5, AR238: 5, AR296: 4, AR184: 4, AR267: 4, AR039: 4, AR275: 4, AR205: 4, AR280: 4, AR246: 4, AR291: 4, AR284: 4, AR286: 4, AR268: 4, AR282: 4, AR281: 4,		

AR289:	4, AR229:	4,							
AR096:	3, AR299:	3,							
AR312:	3, AR298:	3,							
AR089:	3, AR234:	3,							
AR315:	3, AR241:	3,							
AR052:	3, AR253:	3,							
AR290:	3, AR237:	3,							
AR231:	3, AR053:	3,							
AR213:	3, AR033:	3,							
AR249:	3, AR316:	3,							
AR226:	3, AR310:	2,							
AR273:	2, AR232:	2,							
AR060:	2, AR227:	2,							
AR309:	2, AR206:	2,							
AR277:	2, AR104:	2,							
AR247:	2, AR292:	2,							
AR300:	2, AR314:	2,							
AR233:	1, AR177:	1,							
AR240:	1, AR185:	1,							
AR293:	1, AR055:	1,							
AR186:	1, AR283:	1,							
AR175:	1, AR061:	1,							
AR183:	1, AR248:	1,							
AR274:	1, AR198:	1,							
AR244:	1, AR256:	0,							
AR295:	0, AR258:	0,							
AR192:	0, AR259:	0,							
AR179:	0, AR219:	0,							
AR218:	0, AR204:	0,							
AR294:	0								

173	HSIEN53	HSIEN53R	183	2 - 208	359	AR274:1152, AR052:1048, AR267: 969, AR309: 961, AR269: 894, AR273: 803, AR213: 767, AR253: 760, AR271: 722, AR312: 721, AR268: 665, AR179: 648, AR270: 632, AR218: 599, AR182: 582, AR183: 559, AR219: 547, AR290: 540, AR313: 515, AR175: 489, AR249: 456, AR089: 454, AR284: 422, AR060: 411, AR293: 406, AR192: 395, AR240: 385, AR248: 380, AR184: 377, AR185: 369, AR096: 359, AR299: 346, AR316: 331, AR247: 329, AR310: 328, AR289: 323, AR275: 319, AR265: 318, AR280: 317, AR291: 308, AR300: 302, AR053: 301, AR282: 301,			
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174	HWBEA69	HWBEA69	184	220 - 375	360	Gln-1 to Trp-9,	AR292: 289, AR314: 282, AR177: 270, AR186: 265, AR315: 244, AR296: 241, AR266: 230, AR256: 228, AR237: 224, AR198: 217, AR298: 213, AR294: 208, AR251: 206, AR039: 205, AR243: 202, AR231: 197, AR285: 193, AR226: 180, AR232: 172, AR277: 162, AR263: 162, AR104: 156, AR234: 145, AR241: 144, AR238: 143, AR295: 142, AR205: 125, AR229: 119, AR033: 108, AR061: 106, AR259: 105, AR244: 101, AR281: 98, AR286: 78, AR258: 76, AR227: 74, AR233: 71, AR206: 50, AR204: 48, AR246: 46, AR283: 39, AR055: 34, AR194: 20, AR202: 15		
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least 80%, at least 75%, at least 70%, at least 60%, or at least 50% of the activity in absence of the antibody.

[0212] The invention also features receptor-specific antibodies which both prevent ligand binding and receptor activation as well as antibodies that recognize the receptor-ligand complex, and, preferably, do not specifically recognize the unbound receptor or the unbound ligand. Likewise, included in the invention are neutralizing antibodies which bind the ligand and prevent binding of the ligand to the receptor, as well as antibodies which bind the ligand, thereby preventing receptor activation, but do not prevent the ligand from binding the receptor. Further included in the invention are antibodies which activate the receptor. These antibodies may act as receptor agonists, i.e., potentiate or activate either all or a subset of the biological activities of the ligand-mediated receptor activation, for example, by inducing dimerization of the receptor. The antibodies may be specified as agonists, antagonists or inverse agonists for biological activities comprising the specific biological activities of the peptides of the invention disclosed herein. The above antibody agonists can be made using methods known in the art. See, e.g., PCT publication WO 96/40281; U.S. Patent No. 5,811,097; Deng et al., *Blood* 92(6):1981-1988 (1998); Chen et al., *Cancer Res.* 58(16):3668-3678 (1998); Harrop et al., *J. Immunol.* 161(4):1786-1794 (1998); Zhu et al., *Cancer Res.* 58(15):3209-3214 (1998); Yoon et al., *J. Immunol.* 160(7):3170-3179 (1998); Prat et al., *J. Cell. Sci.* 111(Pt2):237-247 (1998); Pitard et al., *J. Immunol. Methods* 205(2):177-190 (1997); Liautard et al., *Cytokine* 9(4):233-241 (1997); Carlson et al., *J. Biol. Chem.* 272(17):11295-11301 (1997); Taryman et al., *Neuron* 14(4):755-762 (1995); Muller et al., *Structure* 6(9):1153-1167 (1998); Bartunek et al., *Cytokine* 8(1):14-20 (1996) (which are all incorporated by reference herein in their entireties).

[0213] Antibodies of the present invention may be used, for example, to purify, detect, and target the polypeptides of the present invention, including both *in vitro* and *in vivo* diagnostic and therapeutic methods. For example, the antibodies have utility in immunoassays for qualitatively and quantitatively measuring levels of the polypeptides of the present invention in biological samples. See, e.g., Harlow et al., *Antibodies: A Laboratory Manual*, (Cold Spring Harbor Laboratory Press, 2nd ed. 1988); incorporated by reference herein in its entirety.

present invention are also included in the present invention. In a specific embodiment, the above-described cross-reactivity is with respect to any single specific antigenic or immunogenic polypeptide, or combination(s) of 2, 3, 4, 5, or more of the specific antigenic and/or immunogenic polypeptides disclosed herein. Further included in the present invention are antibodies which bind polypeptides encoded by polynucleotides which hybridize to a polynucleotide of the present invention under stringent hybridization conditions (as described herein). Antibodies of the present invention may also be described or specified in terms of their binding affinity to a polypeptide of the invention. Preferred binding affinities include those with a dissociation constant or K_d less than 5×10^{-2} M, 10^{-2} M, 5×10^{-3} M, 10^{-3} M, 5×10^{-4} M, 10^{-4} M, 5×10^{-5} M, 10^{-5} M, 5×10^{-6} M, 10^{-6} M, 5×10^{-7} M, 10^{-7} M, 5×10^{-8} M, 10^{-8} M, 5×10^{-9} M, 10^{-9} M, 5×10^{-10} M, 10^{-10} M, 5×10^{-11} M, 10^{-11} M, 5×10^{-12} M, 10^{-12} M, 5×10^{-13} M, 10^{-13} M, 5×10^{-14} M, 10^{-14} M, 5×10^{-15} M, or 10^{-15} M.

[0210] The invention also provides antibodies that competitively inhibit binding of an antibody to an epitope of the invention as determined by any method known in the art for determining competitive binding, for example, the immunoassays described herein. In preferred embodiments, the antibody competitively inhibits binding to the epitope by at least 95%, at least 90%, at least 85 %, at least 80%, at least 75%, at least 70%, at least 60%, or at least 50%.

[0211] Antibodies of the present invention may act as agonists or antagonists of the polypeptides of the present invention. For example, the present invention includes antibodies which disrupt the receptor/ligand interactions with the polypeptides of the invention either partially or fully. Preferably, antibodies of the present invention bind an antigenic epitope disclosed herein, or a portion thereof. The invention features both receptor-specific antibodies and ligand-specific antibodies. The invention also features receptor-specific antibodies which do not prevent ligand binding but prevent receptor activation. Receptor activation (i.e., signaling) may be determined by techniques described herein or otherwise known in the art. For example, receptor activation can be determined by detecting the phosphorylation (e.g., tyrosine or serine/threonine) of the receptor or its substrate by immunoprecipitation followed by western blot analysis (for example, as described *supra*). In specific embodiments, antibodies are provided that inhibit ligand activity or receptor activity by at least 95%, at least 90%, at least 85%, at

immunoglobulins, as described infra and, for example in, U.S. Patent No. 5,939,598 by Kucherlapati et al.

[0207] The antibodies of the present invention may be monospecific, bispecific, trispecific or of greater multispecificity. Multispecific antibodies may be specific for different epitopes of a polypeptide of the present invention or may be specific for both a polypeptide of the present invention as well as for a heterologous epitope, such as a heterologous polypeptide or solid support material. See, e.g., PCT publications WO 93/17715; WO 92/08802; WO 91/00360; WO 92/05793; Tutt, et al., J. Immunol. 147:60-69 (1991); U.S. Patent Nos. 4,474,893; 4,714,681; 4,925,648; 5,573,920; 5,601,819; Kostelny et al., J. Immunol. 148:1547-1553 (1992).

[0208] Antibodies of the present invention may be described or specified in terms of the epitope(s) or portion(s) of a polypeptide of the present invention which they recognize or specifically bind. The epitope(s) or polypeptide portion(s) may be specified as described herein, e.g., by N-terminal and C-terminal positions, or by size in contiguous amino acid residues, or listed in the Tables and Figures. Preferred epitopes of the invention include the predicted epitopes shown in column 7 of Table 1, as well as polynucleotides that encode these epitopes. Antibodies which specifically bind any epitope or polypeptide of the present invention may also be excluded. Therefore, the present invention includes antibodies that specifically bind polypeptides of the present invention, and allows for the exclusion of the same.

[0209] Antibodies of the present invention may also be described or specified in terms of their cross-reactivity. Antibodies that do not bind any other analog, ortholog, or homolog of a polypeptide of the present invention are included. Antibodies that bind polypeptides with at least 95%, at least 90%, at least 85%, at least 80%, at least 75%, at least 70%, at least 65%, at least 60%, at least 55%, and at least 50% identity (as calculated using methods known in the art and described herein) to a polypeptide of the present invention are also included in the present invention. In specific embodiments, antibodies of the present invention cross-react with murine, rat and/or rabbit homologs of human proteins and the corresponding epitopes thereof. Antibodies that do not bind polypeptides with less than 95%, less than 90%, less than 85%, less than 80%, less than 75%, less than 70%, less than 65%, less than 60%, less than 55%, and less than 50% identity (as calculated using methods known in the art and described herein) to a polypeptide of the

variant of the invention (e.g., a polypeptide or fragment or variant of the amino acid sequence of SEQ ID NO:Y or a polypeptide encoded by the cDNA contained in Clone ID No:Z, and/or an epitope, of the present invention) as determined by immunoassays well known in the art for assaying specific antibody-antigen binding. Antibodies of the invention include, but are not limited to, polyclonal, monoclonal, multispecific, human, humanized or chimeric antibodies, single chain antibodies, Fab fragments, F(ab') fragments, fragments produced by a Fab expression library, anti-idiotypic (anti-Id) antibodies (including, e.g., anti-Id antibodies to antibodies of the invention), intracellularly-made antibodies (i.e., intrabodies), and epitope-binding fragments of any of the above. The term "antibody," as used herein, refers to immunoglobulin molecules and immunologically active portions of immunoglobulin molecules, i.e., molecules that contain an antigen binding site that immunospecifically binds an antigen. The immunoglobulin molecules of the invention can be of any type (e.g., IgG, IgE, IgM, IgD, IgA and IgY), class (e.g., IgG1, IgG2, IgG3, IgG4, IgA1 and IgA2) or subclass of immunoglobulin molecule. In preferred embodiments, the immunoglobulin molecules of the invention are IgG1. In other preferred embodiments, the immunoglobulin molecules of the invention are IgG4.

[0206] Most preferably the antibodies are human antigen-binding antibody fragments of the present invention and include, but are not limited to, Fab, Fab' and F(ab')₂, Fd, single-chain Fvs (scFv), single-chain antibodies, disulfide-linked Fvs (sdFv) and fragments comprising either a VL or VH domain. Antigen-binding antibody fragments, including single-chain antibodies, may comprise the variable region(s) alone or in combination with the entirety or a portion of the following: hinge region, CH1, CH2, and CH3 domains. Also included in the invention are antigen-binding fragments also comprising any combination of variable region(s) with a hinge region, CH1, CH2, and CH3 domains. The antibodies of the invention may be from any animal origin including birds and mammals. Preferably, the antibodies are human, murine (e.g., mouse and rat), donkey, sheep rabbit, goat, guinea pig, camel, horse, or chicken. As used herein, "human" antibodies include antibodies having the amino acid sequence of a human immunoglobulin and include antibodies isolated from human immunoglobulin libraries or from animals transgenic for one or more human immunoglobulin and that do not express endogenous

invention may be generated using techniques known in the art to form one or more inter-molecule cross-links between the cysteine residues located within the sequence of the polypeptides desired to be contained in the multimer (see, e.g., US Patent Number 5,478,925, which is herein incorporated by reference in its entirety). Further, polypeptides of the invention may be routinely modified by the addition of cysteine or biotin to the C-terminus or N-terminus of the polypeptide and techniques known in the art may be applied to generate multimers containing one or more of these modified polypeptides (see, e.g., US Patent Number 5,478,925, which is herein incorporated by reference in its entirety). Additionally, techniques known in the art may be applied to generate liposomes containing the polypeptide components desired to be contained in the multimer of the invention (see, e.g., US Patent Number 5,478,925, which is herein incorporated by reference in its entirety).

[0204] Alternatively, multimers of the invention may be generated using genetic engineering techniques known in the art. In one embodiment, polypeptides contained in multimers of the invention are produced recombinantly using fusion protein technology described herein or otherwise known in the art (see, e.g., US Patent Number 5,478,925, which is herein incorporated by reference in its entirety). In a specific embodiment, polynucleotides coding for a homodimer of the invention are generated by ligating a polynucleotide sequence encoding a polypeptide of the invention to a sequence encoding a linker polypeptide and then further to a synthetic polynucleotide encoding the translated product of the polypeptide in the reverse orientation from the original C-terminus to the N-terminus (lacking the leader sequence) (see, e.g., US Patent Number 5,478,925, which is herein incorporated by reference in its entirety). In another embodiment, recombinant techniques described herein or otherwise known in the art are applied to generate recombinant polypeptides of the invention which contain a transmembrane domain (or hydrophobic or signal peptide) and which can be incorporated by membrane reconstitution techniques into liposomes (see, e.g., US Patent Number 5,478,925, which is herein incorporated by reference in its entirety).

Antibodies

[0205] Further polypeptides of the invention relate to antibodies and T-cell antigen receptors (TCR) which immunospecifically bind a polypeptide, polypeptide fragment, or

Proteins comprising multiple polypeptides of the invention separated by peptide linkers may be produced using conventional recombinant DNA technology.

[0200] Another method for preparing multimer polypeptides of the invention involves use of polypeptides of the invention fused to a leucine zipper or isoleucine zipper polypeptide sequence. Leucine zipper and isoleucine zipper domains are polypeptides that promote multimerization of the proteins in which they are found. Leucine zippers were originally identified in several DNA-binding proteins (Landschulz et al., Science 240:1759, (1988)), and have since been found in a variety of different proteins. Among the known leucine zippers are naturally occurring peptides and derivatives thereof that dimerize or trimerize. Examples of leucine zipper domains suitable for producing soluble multimeric proteins of the invention are those described in PCT application WO 94/10308, hereby incorporated by reference. Recombinant fusion proteins comprising a polypeptide of the invention fused to a polypeptide sequence that dimerizes or trimerizes in solution are expressed in suitable host cells, and the resulting soluble multimeric fusion protein is recovered from the culture supernatant using techniques known in the art.

[0201] Trimeric polypeptides of the invention may offer the advantage of enhanced biological activity. Preferred leucine zipper moieties and isoleucine moieties are those that preferentially form trimers. One example is a leucine zipper derived from lung surfactant protein D (SPD), as described in Hoppe et al. (FEBS Letters 344:191, (1994)) and in U.S. patent application Ser. No. 08/446,922, hereby incorporated by reference. Other peptides derived from naturally occurring trimeric proteins may be employed in preparing trimeric polypeptides of the invention.

[0202] In another example, proteins of the invention are associated by interactions between Flag® polypeptide sequence contained in fusion proteins of the invention containing Flag® polypeptide sequence. In a further embodiment, proteins of the invention are associated by interactions between heterologous polypeptide sequence contained in Flag® fusion proteins of the invention and anti-Flag® antibody.

[0203] The multimers of the invention may be generated using chemical techniques known in the art. For example, polypeptides desired to be contained in the multimers of the invention may be chemically cross-linked using linker molecules and linker molecule length optimization techniques known in the art (see, e.g., US Patent Number 5,478,925, which is herein incorporated by reference in its entirety). Additionally, multimers of the

a heterodimer, a heterotrimer, or a heterotetramer. In additional embodiments, the heteromeric multimer of the invention is at least a heterodimer, at least a heterotrimer, or at least a heterotetramer.

[0199] Multimers of the invention may be the result of hydrophobic, hydrophilic, ionic and/or covalent associations and/or may be indirectly linked by, for example, liposome formation. Thus, in one embodiment, multimers of the invention, such as, for example, homodimers or homotrimers, are formed when polypeptides of the invention contact one another in solution. In another embodiment, heteromultimers of the invention, such as, for example, heterotrimers or heterotetramers, are formed when polypeptides of the invention contact antibodies to the polypeptides of the invention (including antibodies to the heterologous polypeptide sequence in a fusion protein of the invention) in solution. In other embodiments, multimers of the invention are formed by covalent associations with and/or between the polypeptides of the invention. Such covalent associations may involve one or more amino acid residues contained in the polypeptide sequence (e.g., that recited in SEQ ID NO:Y, encoded by the portion of SEQ ID NO:X as defined in columns 8 and 9 of Table 2, and/or encoded by the cDNA contained in Clone ID NO:Z). In one instance, the covalent associations are cross-linking between cysteine residues located within the polypeptide sequences which interact in the native (i.e., naturally occurring) polypeptide. In another instance, the covalent associations are the consequence of chemical or recombinant manipulation. Alternatively, such covalent associations may involve one or more amino acid residues contained in the heterologous polypeptide sequence in a fusion protein. In one example, covalent associations are between the heterologous sequence contained in a fusion protein of the invention (see, e.g., US Patent Number 5,478,925). In a specific example, the covalent associations are between the heterologous sequence contained in a Fc fusion protein of the invention (as described herein). In another specific example, covalent associations of fusion proteins of the invention are between heterologous polypeptide sequence from another protein that is capable of forming covalently associated multimers, such as for example, osteoprotegerin (see, e.g., International Publication NO: WO 98/49305, the contents of which are herein incorporated by reference in its entirety). In another embodiment, two or more polypeptides of the invention are joined through peptide linkers. Examples include those peptide linkers described in U.S. Pat. No. 5,073,627 (hereby incorporated by reference).

chromatography, affinity chromatography, hydroxylapatite chromatography and lectin chromatography. Most preferably, high performance liquid chromatography ("HPLC") is employed for purification. Well known techniques for refolding protein may be employed to regenerate active conformation when the polypeptide is denatured during isolation and/or purification.

[0196] The polypeptides of the invention may be in monomers or multimers (i.e., dimers, trimers, tetramers and higher multimers). Accordingly, the present invention relates to monomers and multimers of the polypeptides of the invention, their preparation, and compositions (preferably, Therapeutics) containing them. In specific embodiments, the polypeptides of the invention are monomers, dimers, trimers or tetramers. In additional embodiments, the multimers of the invention are at least dimers, at least trimers, or at least tetramers.

[0197] Multimers encompassed by the invention may be homomers or heteromers. As used herein, the term homomer refers to a multimer containing only polypeptides corresponding to a protein of the invention (e.g., the amino acid sequence of SEQ ID NO:Y, an amino acid sequence encoded by SEQ ID NO:X or the complement of SEQ ID NO:X, the amino acid sequence encoded by the portion of SEQ ID NO:X as defined in columns 8 and 9 of Table 2, and/or an amino acid sequence encoded by cDNA contained in Clone ID NO:Z (including fragments, variants, splice variants, and fusion proteins, corresponding to these as described herein)). These homomers may contain polypeptides having identical or different amino acid sequences. In a specific embodiment, a homomer of the invention is a multimer containing only polypeptides having an identical amino acid sequence. In another specific embodiment, a homomer of the invention is a multimer containing polypeptides having different amino acid sequences. In specific embodiments, the multimer of the invention is a homodimer (e.g., containing two polypeptides having identical or different amino acid sequences) or a homotrimer (e.g., containing three polypeptides having identical and/or different amino acid sequences). In additional embodiments, the homomeric multimer of the invention is at least a homodimer, at least a homotrimer, or at least a homotetramer.

[0198] As used herein, the term heteromer refers to a multimer containing one or more heterologous polypeptides (i.e., polypeptides of different proteins) in addition to the polypeptides of the invention. In a specific embodiment, the multimer of the invention is

[0192] One system for attaching polyethylene glycol directly to amino acid residues of proteins without an intervening linker employs tresylated MPEG, which is produced by the modification of monmethoxy polyethylene glycol (MPEG) using tresylchloride ($\text{ClSO}_2\text{CH}_2\text{CF}_3$). Upon reaction of protein with tresylated MPEG, polyethylene glycol is directly attached to amine groups of the protein. Thus, the invention includes protein-polyethylene glycol conjugates produced by reacting proteins of the invention with a polyethylene glycol molecule having a 2,2,2-trifluoroethane sulphonyl group.

[0193] Polyethylene glycol can also be attached to proteins using a number of different intervening linkers. For example, U.S. Patent No. 5,612,460, the entire disclosure of which is incorporated herein by reference, discloses urethane linkers for connecting polyethylene glycol to proteins. Protein-polyethylene glycol conjugates wherein the polyethylene glycol is attached to the protein by a linker can also be produced by reaction of proteins with compounds such as MPEG-succinimidylsuccinate, MPEG activated with 1,1'-carbonyldiimidazole, MPEG-2,4,5-trichloropenylcarbonate, MPEG-p-nitrophenolcarbonate, and various MPEG-succinate derivatives. A number of additional polyethylene glycol derivatives and reaction chemistries for attaching polyethylene glycol to proteins are described in International Publication No. WO 98/32466, the entire disclosure of which is incorporated herein by reference. Pegylated protein products produced using the reaction chemistries set out herein are included within the scope of the invention.

[0194] The number of polyethylene glycol moieties attached to each protein of the invention (i.e., the degree of substitution) may also vary. For example, the pegylated proteins of the invention may be linked, on average, to 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 12, 15, 17, 20, or more polyethylene glycol molecules. Similarly, the average degree of substitution within ranges such as 1-3, 2-4, 3-5, 4-6, 5-7, 6-8, 7-9, 8-10, 9-11, 10-12, 11-13, 12-14, 13-15, 14-16, 15-17, 16-18, 17-19, or 18-20 polyethylene glycol moieties per protein molecule. Methods for determining the degree of substitution are discussed, for example, in Delgado et al., Crit. Rev. Thera. Drug Carrier Sys. 9:249-304 (1992).

[0195] The polypeptides of the invention can be recovered and purified from chemical synthesis and recombinant cell cultures by standard methods which include, but are not limited to, ammonium sulfate or ethanol precipitation, acid extraction, anion or cation exchange chromatography, phosphocellulose chromatography, hydrophobic interaction

and the C-terminal amino acid residue. Sulfhydryl groups may also be used as a reactive group for attaching the polyethylene glycol molecules. Preferred for therapeutic purposes is attachment at an amino group, such as attachment at the N-terminus or lysine group.

[0189] As suggested above, polyethylene glycol may be attached to proteins via linkage to any of a number of amino acid residues. For example, polyethylene glycol can be linked to proteins via covalent bonds to lysine, histidine, aspartic acid, glutamic acid, or cysteine residues. One or more reaction chemistries may be employed to attach polyethylene glycol to specific amino acid residues (e.g., lysine, histidine, aspartic acid, glutamic acid, or cysteine) of the protein or to more than one type of amino acid residue (e.g., lysine, histidine, aspartic acid, glutamic acid, cysteine and combinations thereof) of the protein.

[0190] One may specifically desire proteins chemically modified at the N-terminus. Using polyethylene glycol as an illustration of the present composition, one may select from a variety of polyethylene glycol molecules (by molecular weight, branching, etc.), the proportion of polyethylene glycol molecules to protein (polypeptide) molecules in the reaction mix, the type of pegylation reaction to be performed, and the method of obtaining the selected N-terminally pegylated protein. The method of obtaining the N-terminally pegylated preparation (i.e., separating this moiety from other monopegylated moieties if necessary) may be by purification of the N-terminally pegylated material from a population of pegylated protein molecules. Selective proteins chemically modified at the N-terminus modification may be accomplished by reductive alkylation which exploits differential reactivity of different types of primary amino groups (lysine versus the N-terminal) available for derivatization in a particular protein. Under the appropriate reaction conditions, substantially selective derivatization of the protein at the N-terminus with a carbonyl group containing polymer is achieved.

[0191] As indicated above, pegylation of the proteins of the invention may be accomplished by any number of means. For example, polyethylene glycol may be attached to the protein either directly or by an intervening linker. Linkerless systems for attaching polyethylene glycol to proteins are described in Delgado et al., *Crit. Rev. Thera. Drug Carrier Sys.* 9:249-304 (1992); Francis et al., *Intern. J. of Hematol.* 68:1-18 (1998); U.S. Patent No. 4,002,531; U.S. Patent No. 5,349,052; WO 95/06058; and WO 98/32466, the disclosures of each of which are incorporated herein by reference.

predetermined positions within the molecule and may include one, two, three or more attached chemical moieties.

[0186] The polymer may be of any molecular weight, and may be branched or unbranched. For polyethylene glycol, the preferred molecular weight is between about 1 kDa and about 100 kDa (the term "about" indicating that in preparations of polyethylene glycol, some molecules will weigh more, some less, than the stated molecular weight) for ease in handling and manufacturing. Other sizes may be used, depending on the desired therapeutic profile (e.g., the duration of sustained release desired, the effects, if any on biological activity, the ease in handling, the degree or lack of antigenicity and other known effects of the polyethylene glycol to a therapeutic protein or analog). For example, the polyethylene glycol may have an average molecular weight of about 200, 500, 1000, 1500, 2000, 2500, 3000, 3500, 4000, 4500, 5000, 5500, 6000, 6500, 7000, 7500, 8000, 8500, 9000, 9500, 10,000, 10,500, 11,000, 11,500, 12,000, 12,500, 13,000, 13,500, 14,000, 14,500, 15,000, 15,500, 16,000, 16,500, 17,000, 17,500, 18,000, 18,500, 19,000, 19,500, 20,000, 25,000, 30,000, 35,000, 40,000, 45,000, 50,000, 55,000, 60,000, 65,000, 70,000, 75,000, 80,000, 85,000, 90,000, 95,000, or 100,000 kDa.

[0187] As noted above, the polyethylene glycol may have a branched structure. Branched polyethylene glycols are described, for example, in U.S. Patent No. 5,643,575; Morpurgo *et al.*, *Appl. Biochem. Biotechnol.* 56:59-72 (1996); Vorobjev *et al.*, *Nucleosides Nucleotides* 18:2745-2750 (1999); and Caliceti *et al.*, *Bioconjug. Chem.* 10:638-646 (1999), the disclosures of each of which are incorporated herein by reference.

[0188] The polyethylene glycol molecules (or other chemical moieties) should be attached to the protein with consideration of effects on functional or antigenic domains of the protein. There are a number of attachment methods available to those skilled in the art, such as, for example, the method disclosed in EP 0 401 384 (coupling PEG to G-CSF), herein incorporated by reference; see also Malik *et al.*, *Exp. Hematol.* 20:1028-1035 (1992), reporting pegylation of GM-CSF using tresyl chloride. For example, polyethylene glycol may be covalently bound through amino acid residues via a reactive group, such as a free amino or carboxyl group. Reactive groups are those to which an activated polyethylene glycol molecule may be bound. The amino acid residues having a free amino group may include lysine residues and the N-terminal amino acid residues; those having a free carboxyl group may include aspartic acid residues glutamic acid residues

Cancer Res. 4(10):2483-90 (1998); Peterson et al., Bioconjug. Chem. 10(4):553-7 (1999); and Zimmerman et al, Nucl. Med. Biol. 26(8):943-50 (1999); which are hereby incorporated by reference in their entirety.

[0184] As mentioned, the proteins of the invention may be modified by either natural processes, such as posttranslational processing, or by chemical modification techniques which are well known in the art. It will be appreciated that the same type of modification may be present in the same or varying degrees at several sites in a given polypeptide. Polypeptides of the invention may be branched, for example, as a result of ubiquitination, and they may be cyclic, with or without branching. Cyclic, branched, and branched cyclic polypeptides may result from posttranslation natural processes or may be made by synthetic methods. Modifications include acetylation, acylation, ADP-ribosylation, amidation, covalent attachment of flavin, covalent attachment of a heme moiety, covalent attachment of a nucleotide or nucleotide derivative, covalent attachment of a lipid or lipid derivative, covalent attachment of phosphatidylinositol, cross-linking, cyclization, disulfide bond formation, demethylation, formation of covalent cross-links, formation of cysteine, formation of pyroglutamate, formylation, gamma-carboxylation, glycosylation, GPI anchor formation, hydroxylation, iodination, methylation, myristoylation, oxidation, pegylation, proteolytic processing, phosphorylation, prenylation, racemization, selenoylation, sulfation, transfer-RNA mediated addition of amino acids to proteins such as arginylation, and ubiquitination. (See, for instance, PROTEINS - STRUCTURE AND MOLECULAR PROPERTIES, 2nd Ed., T. E. Creighton, W. H. Freeman and Company, New York (1993); POSTTRANSLATIONAL COVALENT MODIFICATION OF PROTEINS, B. C. Johnson, Ed., Academic Press, New York, pgs. 1-12 (1983); Seifter et al., Meth. Enzymol. 182:626-646 (1990); Rattan et al., Ann. N.Y. Acad. Sci. 663:48-62 (1992)).

[0185] Also provided by the invention are chemically modified derivatives of the polypeptides of the invention which may provide additional advantages such as increased solubility, stability and circulating time of the polypeptide, or decreased immunogenicity (see U.S. Patent No. 4,179,337). The chemical moieties for derivitization may be selected from water soluble polymers such as polyethylene glycol, ethylene glycol/propylene glycol copolymers, carboxymethylcellulose, dextran, polyvinyl alcohol and the like. The polypeptides may be modified at random positions within the molecule, or at

not limited, to specific chemical cleavage by cyanogen bromide, trypsin, chymotrypsin, papain, V8 protease, NaBH₄; acetylation, formylation, oxidation, reduction; metabolic synthesis in the presence of tunicamycin; etc.

[0181] Additional post-translational modifications encompassed by the invention include, for example, e.g., N-linked or O-linked carbohydrate chains, processing of N-terminal or C-terminal ends), attachment of chemical moieties to the amino acid backbone, chemical modifications of N-linked or O-linked carbohydrate chains, and addition or deletion of an N-terminal methionine residue as a result of procaryotic host cell expression. The polypeptides may also be modified with a detectable label, such as an enzymatic, fluorescent, isotopic or affinity label to allow for detection and isolation of the protein.

[0182] Examples of suitable enzymes include horseradish peroxidase, alkaline phosphatase, beta-galactosidase, or acetylcholinesterase; examples of suitable prosthetic group complexes include streptavidin/biotin and avidin/biotin; examples of suitable fluorescent materials include umbelliferone, fluorescein, fluorescein isothiocyanate, rhodamine, dichlorotriazinylamine fluorescein, dansyl chloride or phycoerythrin; an example of a luminescent material includes luminol; examples of bioluminescent materials include luciferase, luciferin, and aequorin; and examples of suitable radioactive material include iodine (¹²¹I, ¹²³I, ¹²⁵I, ¹³¹I), carbon (¹⁴C), sulfur (³⁵S), tritium (³H), indium (¹¹¹In, ¹¹²In, ^{113m}In, ^{115m}In), technetium (⁹⁹Tc, ^{99m}Tc), thallium (²⁰¹Tl), gallium (⁶⁸Ga, ⁶⁷Ga), palladium (¹⁰³Pd), molybdenum (⁹⁹Mo), xenon (¹³³Xe), fluorine (¹⁸F), ¹⁵³Sm, ¹⁷⁷Lu, ¹⁵⁹Gd, ¹⁴⁹Pm, ¹⁴⁰La, ¹⁷⁵Yb, ¹⁶⁶Ho, ⁹⁰Y, ⁴⁷Sc, ¹⁸⁶Re, ¹⁸⁸Re, ¹⁴²Pr, ¹⁰⁵Rh, and ⁹⁷Ru.

[0183] In specific embodiments, a polypeptide of the present invention or fragment or variant thereof is attached to macrocyclic chelators that associate with radiometal ions, including but not limited to, ¹⁷⁷Lu, ⁹⁰Y, ¹⁶⁶Ho, and ¹⁵³Sm, to polypeptides. In a preferred embodiment, the radiometal ion associated with the macrocyclic chelators is ¹¹¹In. In another preferred embodiment, the radiometal ion associated with the macrocyclic chelator is ⁹⁰Y. In specific embodiments, the macrocyclic chelator is 1,4,7,10-tetraazacyclododecane-N,N',N'',N'''-tetraacetic acid (DOTA). In other specific embodiments, DOTA is attached to an antibody of the invention or fragment thereof via a linker molecule. Examples of linker molecules useful for conjugating DOTA to a polypeptide are commonly known in the art - see, for example, DeNardo et al., Clin

host cells of vertebrate origin, particularly mammalian origin, that have been engineered to delete or replace endogenous genetic material (e.g., coding sequence), and/or to include genetic material (e.g., heterologous polynucleotide sequences) that is operably associated with polynucleotides of the invention, and which activates, alters, and/or amplifies endogenous polynucleotides. For example, techniques known in the art may be used to operably associate heterologous control regions (e.g., promoter and/or enhancer) and endogenous polynucleotide sequences via homologous recombination (see, e.g., U.S. Patent No. 5,641,670, issued June 24, 1997; International Publication No. WO 96/29411, published September 26, 1996; International Publication No. WO 94/12650, published August 4, 1994; Koller et al., Proc. Natl. Acad. Sci. USA 86:8932-8935 (1989); and Zijlstra et al., Nature 342:435-438 (1989), the disclosures of each of which are incorporated by reference in their entireties).

[0179] In addition, polypeptides of the invention can be chemically synthesized using techniques known in the art (e.g., see Creighton, 1983, *Proteins: Structures and Molecular Principles*, W.H. Freeman & Co., N.Y., and Hunkapiller et al., *Nature*, 310:105-111 (1984)). For example, a polypeptide corresponding to a fragment of a polypeptide can be synthesized by use of a peptide synthesizer. Furthermore, if desired, nonclassical amino acids or chemical amino acid analogs can be introduced as a substitution or addition into the polypeptide sequence. Non-classical amino acids include, but are not limited to, to the D-isomers of the common amino acids, 2,4-diaminobutyric acid, α -amino isobutyric acid, 4-aminobutyric acid, Abu, 2-amino butyric acid, g-Abu, e-Ahx, 6-amino hexanoic acid, Aib, 2-amino isobutyric acid, 3-amino propionic acid, ornithine, norleucine, norvaline, hydroxyproline, sarcosine, citrulline, homocitrulline, cysteic acid, t-butylglycine, t-butylalanine, phenylglycine, cyclohexylalanine, b-alanine, fluoro-amino acids, designer amino acids such as b-methyl amino acids, Ca-methyl amino acids, Na-methyl amino acids, and amino acid analogs in general. Furthermore, the amino acid can be D (dextrorotary) or L (levorotary).

[0180] The invention encompasses polypeptides of the present invention which are differentially modified during or after translation, e.g., by glycosylation, acetylation, phosphorylation, amidation, derivatization by known protecting/blocking groups, proteolytic cleavage, linkage to an antibody molecule or other cellular ligand, etc. Any of numerous chemical modifications may be carried out by known techniques, including but

its sole carbon source, *Pichia pastoris* must generate high levels of alcohol oxidase due, in part, to the relatively low affinity of alcohol oxidase for O₂. Consequently, in a growth medium depending on methanol as a main carbon source, the promoter region of one of the two alcohol oxidase genes (*AOX1*) is highly active. In the presence of methanol, alcohol oxidase produced from the *AOX1* gene comprises up to approximately 30% of the total soluble protein in *Pichia pastoris*. See Ellis, S.B., *et al.*, *Mol. Cell. Biol.* 5:1111-21 (1985); Koutz, P.J., *et al.*, *Yeast* 5:167-77 (1989); Tschopp, J.F., *et al.*, *Nucl. Acids Res.* 15:3859-76 (1987). Thus, a heterologous coding sequence, such as, for example, a polynucleotide of the present invention, under the transcriptional regulation of all or part of the *AOX1* regulatory sequence is expressed at exceptionally high levels in *Pichia* yeast grown in the presence of methanol.

[0175] In one example, the plasmid vector pPIC9K is used to express DNA encoding a polypeptide of the invention, as set forth herein, in a *Pichea* yeast system essentially as described in "*Pichia* Protocols: Methods in Molecular Biology," D.R. Higgins and J. Cregg, eds. The Humana Press, Totowa, NJ, 1998. This expression vector allows expression and secretion of a polypeptide of the invention by virtue of the strong *AOX1* promoter linked to the *Pichia pastoris* alkaline phosphatase (PHO) secretory signal peptide (i.e., leader) located upstream of a multiple cloning site.

[0176] Many other yeast vectors could be used in place of pPIC9K, such as, pYES2, pYD1, pTEF1/Zeo, pYES2/GS, pPICZ, pGAPZ, pGAPZalpha, pPIC9, pPIC3.5, pHIL-D2, pHIL-S1, pPIC3.5K, and PAO815, as one skilled in the art would readily appreciate, as long as the proposed expression construct provides appropriately located signals for transcription, translation, secretion (if desired), and the like, including an in-frame AUG as required.

[0177] In another embodiment, high-level expression of a heterologous coding sequence, such as, for example, a polynucleotide of the present invention, may be achieved by cloning the heterologous polynucleotide of the invention into an expression vector such as, for example, pGAPZ or pGAPZalpha, and growing the yeast culture in the absence of methanol.

[0178] In addition to encompassing host cells containing the vector constructs discussed herein, the invention also encompasses primary, secondary, and immortalized

amplifies endogenous polynucleotides. For example, techniques known in the art may be used to operably associate heterologous control regions (e.g., promoter and/or enhancer) and endogenous polynucleotide sequences via homologous recombination (see, e.g., US Patent Number 5,641,670, issued June 24, 1997; International Publication Number WO 96/29411; International Publication Number WO 94/12650; Koller *et al.*, *Proc. Natl. Acad. Sci. USA* 86:8932-8935 (1989); and Zijlstra *et al.*, *Nature* 342:435-438 (1989), the disclosures of each of which are incorporated by reference in their entireties).

[0172] Polypeptides of the invention can be recovered and purified from recombinant cell cultures by well-known methods including ammonium sulfate or ethanol precipitation, acid extraction, anion or cation exchange chromatography, phosphocellulose chromatography, hydrophobic interaction chromatography, affinity chromatography, hydroxylapatite chromatography and lectin chromatography. Most preferably, high performance liquid chromatography ("HPLC") is employed for purification.

[0173] Polypeptides of the present invention can also be recovered from: products purified from natural sources, including bodily fluids, tissues and cells, whether directly isolated or cultured; products of chemical synthetic procedures; and products produced by recombinant techniques from a prokaryotic or eukaryotic host, including, for example, bacterial, yeast, higher plant, insect, and mammalian cells. Depending upon the host employed in a recombinant production procedure, the polypeptides of the present invention may be glycosylated or may be non-glycosylated. In addition, polypeptides of the invention may also include an initial modified methionine residue, in some cases as a result of host-mediated processes. Thus, it is well known in the art that the N-terminal methionine encoded by the translation initiation codon generally is removed with high efficiency from any protein after translation in all eukaryotic cells. While the N-terminal methionine on most proteins also is efficiently removed in most prokaryotes, for some proteins, this prokaryotic removal process is inefficient, depending on the nature of the amino acid to which the N-terminal methionine is covalently linked.

[0174] In one embodiment, the yeast *Pichia pastoris* is used to express polypeptides of the invention in a eukaryotic system. *Pichia pastoris* is a methylotrophic yeast which can metabolize methanol as its sole carbon source. A main step in the methanol metabolism pathway is the oxidation of methanol to formaldehyde using O₂. This reaction is catalyzed by the enzyme alcohol oxidase. In order to metabolize methanol as

synthase expression vectors can be obtained from Lonza Biologics, Inc. (Portsmouth, NH). Expression and production of monoclonal antibodies using a GS expression system in murine myeloma cells is described in Bebbington *et al.*, *Bio/technology* 10:169(1992) and in Biblia and Robinson *Biotechnol. Prog.* 11:1 (1995) which are herein incorporated by reference.

[0169] The present invention also relates to host cells containing the above-described vector constructs described herein, and additionally encompasses host cells containing nucleotide sequences of the invention that are operably associated with one or more heterologous control regions (e.g., promoter and/or enhancer) using techniques known of in the art. The host cell can be a higher eukaryotic cell, such as a mammalian cell (e.g., a human derived cell), or a lower eukaryotic cell, such as a yeast cell, or the host cell can be a prokaryotic cell, such as a bacterial cell. A host strain may be chosen which modulates the expression of the inserted gene sequences, or modifies and processes the gene product in the specific fashion desired. Expression from certain promoters can be elevated in the presence of certain inducers; thus expression of the genetically engineered polypeptide may be controlled. Furthermore, different host cells have characteristics and specific mechanisms for the translational and post-translational processing and modification (e.g., phosphorylation, cleavage) of proteins. Appropriate cell lines can be chosen to ensure the desired modifications and processing of the foreign protein expressed.

[0170] Introduction of the nucleic acids and nucleic acid constructs of the invention into the host cell can be effected by calcium phosphate transfection, DEAE-dextran mediated transfection, cationic lipid-mediated transfection, electroporation, transduction, infection, or other methods. Such methods are described in many standard laboratory manuals, such as Davis *et al.*, *Basic Methods In Molecular Biology* (1986). It is specifically contemplated that the polypeptides of the present invention may in fact be expressed by a host cell lacking a recombinant vector.

[0171] In addition to encompassing host cells containing the vector constructs discussed herein, the invention also encompasses primary, secondary, and immortalized host cells of vertebrate origin, particularly mammalian origin, that have been engineered to delete or replace endogenous genetic material (e.g., the coding sequence), and/or to include genetic material (e.g., heterologous polynucleotide sequences) that is operably associated with polynucleotides of the invention, and which activates, alters, and/or

codon at the beginning and a termination codon (UAA, UGA or UAG) appropriately positioned at the end of the polypeptide to be translated.

[0166] As indicated, the expression vectors will preferably include at least one selectable marker. Such markers include dihydrofolate reductase, G418, glutamine synthase, or neomycin resistance for eukaryotic cell culture, and tetracycline, kanamycin or ampicillin resistance genes for culturing in *E. coli* and other bacteria. Representative examples of appropriate hosts include, but are not limited to, bacterial cells, such as *E. coli*, *Streptomyces* and *Salmonella typhimurium* cells; fungal cells, such as yeast cells (e.g., *Saccharomyces cerevisiae* or *Pichia pastoris* (ATCC Accession No. 201178)); insect cells such as *Drosophila* S2 and *Spodoptera Sf9* cells; animal cells such as CHO, COS, 293, and Bowes melanoma cells; and plant cells. Appropriate culture mediums and conditions for the above-described host cells are known in the art.

[0167] Among vectors preferred for use in bacteria include pQE70, pQE60 and pQE-9, available from QIAGEN, Inc.; pBluescript vectors, Phagescript vectors, pNH8A, pNH16a, pNH18A, pNH46A, available from Stratagene Cloning Systems, Inc.; and ptrc99a, pKK223-3, pKK233-3, pDR540, pRIT5 available from Pharmacia Biotech, Inc. Among preferred eukaryotic vectors are pWLNEO, pSV2CAT, pOG44, pXT1 and pSG available from Stratagene; and pSVK3, pBPV, pMSG and pSVL available from Pharmacia. Preferred expression vectors for use in yeast systems include, but are not limited to pYES2, pYD1, pTEF1/Zeo, pYES2/GS, pPICZ, pGAPZ, pGAPZalph, pPIC9, pPIC3.5, pHIL-D2, pHIL-S1, pPIC3.5K, pPIC9K, and PAO815 (all available from Invitrogen, Carlsbad, CA). Other suitable vectors will be readily apparent to the skilled artisan.

[0168] Vectors which use glutamine synthase (GS) or DHFR as the selectable markers can be amplified in the presence of the drugs methionine sulfoximine or methotrexate, respectively. An advantage of glutamine synthase based vectors are the availability of cell lines (e.g., the murine myeloma cell line, NS0) which are glutamine synthase negative. Glutamine synthase expression systems can also function in glutamine synthase expressing cells (e.g., Chinese Hamster Ovary (CHO) cells) by providing additional inhibitor to prevent the functioning of the endogenous gene. A glutamine synthase expression system and components thereof are detailed in PCT publications: WO87/04462; WO86/05807; WO89/01036; WO89/10404; and WO91/06657, which are hereby incorporated in their entireties by reference herein. Additionally, glutamine

embodiment, alteration of polynucleotides corresponding to SEQ ID NO:X and the polypeptides encoded by these polynucleotides may be achieved by DNA shuffling. DNA shuffling involves the assembly of two or more DNA segments by homologous or site-specific recombination to generate variation in the polynucleotide sequence. In another embodiment, polynucleotides of the invention, or the encoded polypeptides, may be altered by being subjected to random mutagenesis by error-prone PCR, random nucleotide insertion or other methods prior to recombination. In another embodiment, one or more components, motifs, sections, parts, domains, fragments, etc., of a polynucleotide encoding a polypeptide of the invention may be recombined with one or more components, motifs, sections, parts, domains, fragments, etc. of one or more heterologous molecules.

[0162] Thus, any of these above fusions can be engineered using the polynucleotides or the polypeptides of the present invention.

Recombinant and Synthetic Production of Polypeptides of the Invention

[0163] The present invention also relates to vectors containing the polynucleotide of the present invention, host cells, and the production of polypeptides by synthetic and recombinant techniques. The vector may be, for example, a phage, plasmid, viral, or retroviral vector. Retroviral vectors may be replication competent or replication defective. In the latter case, viral propagation generally will occur only in complementing host cells.

[0164] The polynucleotides of the invention may be joined to a vector containing a selectable marker for propagation in a host. Generally, a plasmid vector is introduced in a precipitate, such as a calcium phosphate precipitate, or in a complex with a charged lipid. If the vector is a virus, it may be packaged in vitro using an appropriate packaging cell line and then transduced into host cells.

[0165] The polynucleotide insert should be operatively linked to an appropriate promoter, such as the phage lambda PL promoter, the E. coli lac, trp, phoA and tac promoters, the SV40 early and late promoters and promoters of retroviral LTRs, to name a few. Other suitable promoters will be known to the skilled artisan. The expression constructs will further contain sites for transcription initiation, termination, and, in the transcribed region, a ribosome binding site for translation. The coding portion of the transcripts expressed by the constructs will preferably include a translation initiating

413 622, and U.S. Patent No. 5,766,883, issued June 16, 1998, herein incorporated by reference in their entirety)), resulting in chimeric polypeptides. For example, EP-A-O 464 533 (Canadian counterpart 2045869) discloses fusion proteins comprising various portions of constant region of immunoglobulin molecules together with another human protein or part thereof. In many cases, the Fc part in a fusion protein is beneficial in therapy and diagnosis, and thus can result in, for example, improved pharmacokinetic properties (EP-A 0232 262). Alternatively, deleting the Fc part after the fusion protein has been expressed, detected, and purified, would be desired. For example, the Fc portion may hinder therapy and diagnosis if the fusion protein is used as an antigen for immunizations. In drug discovery, for example, human proteins, such as hIL-5, have been fused with Fc portions for the purpose of high-throughput screening assays to identify antagonists of hIL-5. See, D. Bennett et al., *J. Molecular Recognition* 8:52-58 (1995); K. Johanson et al., *J. Biol. Chem.* 270:9459-9471 (1995).

[0160] Moreover, the polypeptides of the present invention can be fused to marker sequences, such as a polypeptide which facilitates purification of the fused polypeptide. In preferred embodiments, the marker amino acid sequence is a hexa-histidine peptide, such as the tag provided in a pQE vector (QIAGEN, Inc., 9259 Eton Avenue, Chatsworth, CA, 91311), among others, many of which are commercially available. As described in Gentz et al., *Proc. Natl. Acad. Sci. USA* 86:821-824 (1989), for instance, hexa-histidine provides for convenient purification of the fusion protein. Another peptide tag useful for purification, the "HA" tag, corresponds to an epitope derived from the influenza hemagglutinin protein (Wilson et al., *Cell* 37:767 (1984)).

[0161] Additional fusion proteins of the invention may be generated through the techniques of gene-shuffling, motif-shuffling, exon-shuffling, and/or codon-shuffling (collectively referred to as "DNA shuffling"). DNA shuffling may be employed to modulate the activities of polypeptides of the invention, such methods can be used to generate polypeptides with altered activity, as well as agonists and antagonists of the polypeptides. See, generally, U.S. Patent Nos. 5,605,793; 5,811,238; 5,830,721; 5,834,252; and 5,837,458, and Patten et al., *Curr. Opinion Biotechnol.* 8:724-33 (1997); Harayama, *Trends Biotechnol.* 16(2):76-82 (1998); Hansson, et al., *J. Mol. Biol.* 287:265-76 (1999); and Lorenzo and Blasco, *Biotechniques* 24(2):308-13 (1998) (each of these patents and publications are hereby incorporated by reference in its entirety). In one

protein, can be used as an antigenic tag. Antibodies raised against the polypeptide of the present invention can be used to indirectly detect the second protein by binding to the polypeptide. Moreover, because secreted proteins target cellular locations based on trafficking signals, polypeptides of the present invention which are shown to be secreted can be used as targeting molecules once fused to other proteins.

[0156] Examples of domains that can be fused to polypeptides of the present invention include not only heterologous signal sequences, but also other heterologous functional regions. The fusion does not necessarily need to be direct, but may occur through linker sequences.

[0157] In certain preferred embodiments, proteins of the invention are fusion proteins comprising an amino acid sequence that is an N and/or C- terminal deletion of a polypeptide of the invention. In preferred embodiments, the invention is directed to a fusion protein comprising an amino acid sequence that is at least 90%, 95%, 96%, 97%, 98% or 99% identical to a polypeptide sequence of the invention. Polynucleotides encoding these proteins are also encompassed by the invention.

[0158] Moreover, fusion proteins may also be engineered to improve characteristics of the polypeptide of the present invention. For instance, a region of additional amino acids, particularly charged amino acids, may be added to the N-terminus of the polypeptide to improve stability and persistence during purification from the host cell or subsequent handling and storage. Also, peptide moieties may be added to the polypeptide to facilitate purification. Such regions may be removed prior to final preparation of the polypeptide. The addition of peptide moieties to facilitate handling of polypeptides are familiar and routine techniques in the art.

[0159] As one of skill in the art will appreciate that, as discussed above, polypeptides of the present invention, and epitope-bearing fragments thereof, can be combined with heterologous polypeptide sequences. For example, the polypeptides of the present invention may be fused with heterologous polypeptide sequences, for example, the polypeptides of the present invention may be fused with the constant domain of immunoglobulins (IgA, IgE, IgG, IgM) or portions thereof (CH1, CH2, CH3, and any combination thereof, including both entire domains and portions thereof), or albumin (including, but not limited to, native or recombinant human albumin or fragments or variants thereof (see, e.g., U.S. Patent No. 5,876,969, issued March 2, 1999, EP Patent 0

albumin, where z is an integer from 369 to 419, as described in U.S. Patent 5,766,883 herein incorporated by reference in its entirety. Polypeptides and/or antibodies of the present invention (including fragments or variants thereof) may be fused to either the N- or C-terminal end of the heterologous protein (e.g., immunoglobulin Fc polypeptide or human serum albumin polypeptide). Polynucleotides encoding fusion proteins of the invention are also encompassed by the invention.

[0154] Such fusion proteins as those described above may facilitate purification and may increase half-life *in vivo*. This has been shown for chimeric proteins consisting of the first two domains of the human CD4-polypeptide and various domains of the constant regions of the heavy or light chains of mammalian immunoglobulins. See, e.g., EP 394,827; Traunecker et al., *Nature*, 331:84-86 (1988). Enhanced delivery of an antigen across the epithelial barrier to the immune system has been demonstrated for antigens (e.g., insulin) conjugated to an FcRn binding partner such as IgG or Fc fragments (see, e.g., PCT Publications WO 96/22024 and WO 99/04813). IgG fusion proteins that have a disulfide-linked dimeric structure due to the IgG portion disulfide bonds have also been found to be more efficient in binding and neutralizing other molecules than monomeric polypeptides or fragments thereof alone. See, e.g., Fountoulakis et al., *J. Biochem.*, 270:3958-3964 (1995). Nucleic acids encoding the above epitopes can also be recombined with a gene of interest as an epitope tag (e.g., the hemagglutinin (HA) tag or flag tag) to aid in detection and purification of the expressed polypeptide. For example, a system described by Janknecht et al. allows for the ready purification of non-denatured fusion proteins expressed in human cell lines (Janknecht et al., 1991, *Proc. Natl. Acad. Sci. USA* 88:8972- 897). In this system, the gene of interest is subcloned into a vaccinia recombination plasmid such that the open reading frame of the gene is translationally fused to an amino-terminal tag consisting of six histidine residues. The tag serves as a matrix binding domain for the fusion protein. Extracts from cells infected with the recombinant vaccinia virus are loaded onto Ni²⁺ nitriloacetic acid-agarose column and histidine-tagged proteins can be selectively eluted with imidazole-containing buffers.

Fusion Proteins

[0155] Any polypeptide of the present invention can be used to generate fusion proteins. For example, the polypeptide of the present invention, when fused to a second

such as keyhole limpet hemacyanin (KLH) or tetanus toxoid. For instance, peptides containing cysteine residues may be coupled to a carrier using a linker such as maleimidobenzoyl- N-hydroxysuccinimide ester (MBS), while other peptides may be coupled to carriers using a more general linking agent such as glutaraldehyde. Animals such as rabbits, rats and mice are immunized with either free or carrier- coupled peptides, for instance, by intraperitoneal and/or intradermal injection of emulsions containing about 100 μ g of peptide or carrier protein and Freund's adjuvant or any other adjuvant known for stimulating an immune response. Several booster injections may be needed, for instance, at intervals of about two weeks, to provide a useful titer of anti-peptide antibody which can be detected, for example, by ELISA assay using free peptide adsorbed to a solid surface. The titer of anti-peptide antibodies in serum from an immunized animal may be increased by selection of anti-peptide antibodies, for instance, by adsorption to the peptide on a solid support and elution of the selected antibodies according to methods well known in the art.

[0153]. As one of skill in the art will appreciate, and as discussed above, the polypeptides of the present invention (e.g., those comprising an immunogenic or antigenic epitope) can be fused to heterologous polypeptide sequences. For example, polypeptides of the present invention (including fragments or variants thereof), may be fused with the constant domain of immunoglobulins (IgA, IgE, IgG, IgM), or portions thereof (CH1, CH2, CH3, or any combination thereof and portions thereof, resulting in chimeric polypeptides. By way of another non-limiting example, polypeptides and/or antibodies of the present invention (including fragments or variants thereof) may be fused with albumin (including but not limited to recombinant human serum albumin or fragments or variants thereof (see, e.g., U.S. Patent No. 5,876,969, issued March 2, 1999, EP Patent 0 413 622, and U.S. Patent No. 5,766,883, issued June 16, 1998, herein incorporated by reference in their entirety)). In a preferred embodiment, polypeptides and/or antibodies of the present invention (including fragments or variants thereof) are fused with the mature form of human serum albumin (i.e., amino acids 1 – 585 of human serum albumin as shown in Figures 1 and 2 of EP Patent 0 322 094) which is herein incorporated by reference in its entirety. In another preferred embodiment, polypeptides and/or antibodies of the present invention (including fragments or variants thereof) are fused with polypeptide fragments comprising, or alternatively consisting of, amino acid residues 1-z of human serum

of, at least one, two, three, four, five, six or more of the portion(s) of SEQ ID NO:Y specified in column 7 of Table 1. These polypeptide fragments have been determined to bear antigenic epitopes of the proteins of the invention by the analysis of the Jameson-Wolf antigenic index which is included in the DNASTar suite of computer programs. By "comprise" it is intended that a polypeptide contains at least one, two, three, four, five, six or more of the portion(s) of SEQ ID NO:Y shown in column 7 of Table 1, but it may contain additional flanking residues on either the amino or carboxyl termini of the recited portion. Such additional flanking sequences are preferably sequences naturally found adjacent to the portion; i.e., contiguous sequence shown in SEQ ID NO:Y. The flanking sequence may, however, be sequences from a heterologous polypeptide, such as from another protein described herein or from a heterologous polypeptide not described herein. In particular embodiments, epitope portions of a polypeptide of the invention comprise one, two, three, or more of the portions of SEQ ID NO:Y shown in column 7 of Table 1.

[0151] Similarly, immunogenic epitopes can be used, for example, to induce antibodies according to methods well known in the art. See, for instance, Sutcliffe et al., *supra*; Wilson et al., *supra*; Chow et al., Proc. Natl. Acad. Sci. USA 82:910-914; and Bittle et al., J. Gen. Virol. 66:2347-2354 (1985). Preferred immunogenic epitopes include the immunogenic epitopes disclosed herein, as well as any combination of two, three, four, five or more of these immunogenic epitopes. The polypeptides comprising one or more immunogenic epitopes may be presented for eliciting an antibody response together with a carrier protein, such as an albumin, to an animal system (such as rabbit or mouse), or, if the polypeptide is of sufficient length (at least about 25 amino acids), the polypeptide may be presented without a carrier. However, immunogenic epitopes comprising as few as 8 to 10 amino acids have been shown to be sufficient to raise antibodies capable of binding to, at the very least, linear epitopes in a denatured polypeptide (e.g., in Western blotting).

[0152] Epitope-bearing polypeptides of the present invention may be used to induce antibodies according to methods well known in the art including, but not limited to, *in vivo* immunization, *in vitro* immunization, and phage display methods. See, e.g., Sutcliffe et al., *supra*; Wilson et al., *supra*, and Bittle et al., J. Gen. Virol., 66:2347-2354 (1985). If *in vivo* immunization is used, animals may be immunized with free peptide; however, anti-peptide antibody titer may be boosted by coupling the peptide to a macromolecular carrier,

[0147] The term "epitopes," as used herein, refers to portions of a polypeptide having antigenic or immunogenic activity in an animal, preferably a mammal, and most preferably in a human. In a preferred embodiment, the present invention encompasses a polypeptide comprising an epitope, as well as the polynucleotide encoding this polypeptide. An "immunogenic epitope," as used herein, is defined as a portion of a protein that elicits an antibody response in an animal, as determined by any method known in the art, for example, by the methods for generating antibodies described *infra*. (See, for example, Geysen et al., Proc. Natl. Acad. Sci. USA 81:3998-4002 (1983)). The term "antigenic epitope," as used herein, is defined as a portion of a protein to which an antibody can immunospecifically bind its antigen as determined by any method well known in the art, for example, by the immunoassays described herein. Immunospecific binding excludes non-specific binding but does not necessarily exclude cross-reactivity with other antigens. Antigenic epitopes need not necessarily be immunogenic.

[0148] Fragments which function as epitopes may be produced by any conventional means. (See, e.g., Houghten, R. A., Proc. Natl. Acad. Sci. USA 82:5131-5135 (1985) further described in U.S. Patent No. 4,631,211.)

[0149] In the present invention, antigenic epitopes preferably contain a sequence of at least 4, at least 5, at least 6, at least 7, more preferably at least 8, at least 9, at least 10, at least 11, at least 12, at least 13, at least 14, at least 15, at least 20, at least 25, at least 30, at least 40, at least 50, and, most preferably, between about 15 to about 30 amino acids. Preferred polypeptides comprising immunogenic or antigenic epitopes are at least 10, 15, 20, 25, 30, 35, 40, 45, 50, 55, 60, 65, 70, 75, 80, 85, 90, 95, or 100 amino acid residues in length. Additional non-exclusive preferred antigenic epitopes include the antigenic epitopes disclosed herein, as well as portions thereof. Antigenic epitopes are useful, for example, to raise antibodies, including monoclonal antibodies, that specifically bind the epitope. Preferred antigenic epitopes include the antigenic epitopes disclosed herein, as well as any combination of two, three, four, five or more of these antigenic epitopes. Antigenic epitopes can be used as the target molecules in immunoassays. (See, for instance, Wilson et al., Cell 37:767-778 (1984); Sutcliffe et al., Science 219:660-666 (1983)).

[0150] Non-limiting examples of epitopes of polypeptides that can be used to generate antibodies of the invention include a polypeptide comprising, or alternatively consisting

biological activity) of the polypeptide sequence of which the amino acid sequence is a fragment. By a polypeptide displaying a "functional activity" is meant a polypeptide capable of one or more known functional activities associated with a full-length protein, such as, for example, biological activity, antigenicity, immunogenicity, and/or multimerization, as described herein.

[0144] Other preferred polypeptide fragments are biologically active fragments. Biologically active fragments are those exhibiting activity similar, but not necessarily identical, to an activity of the polypeptide of the present invention. The biological activity of the fragments may include an improved desired activity, or a decreased undesirable activity.

[0145] In preferred embodiments, polypeptides of the invention comprise, or alternatively consist of, one, two, three, four, five or more of the antigenic fragments of the polypeptide of SEQ ID NO:Y, or portions thereof. Polynucleotides encoding these polypeptides are also encompassed by the invention.

[0146] The present invention encompasses polypeptides comprising, or alternatively consisting of, an epitope of: the polypeptide sequence shown in SEQ ID NO:Y; a polypeptide sequence encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide sequence encoded by the portion of SEQ ID NO:X as defined in columns 8 and 9 of Table 2; the polypeptide sequence encoded by the cDNA contained in Clone ID NO:Z; or the polypeptide sequence encoded by a polynucleotide that hybridizes to the sequence of SEQ ID NO:X, the complement of the sequence of SEQ ID NO:X, the complement of a portion of SEQ ID NO:X as defined in columns 8 and 9 of Table 2, or the cDNA sequence contained in Clone ID NO:Z under stringent hybridization conditions or alternatively, under lower stringency hybridization as defined *supra*. The present invention further encompasses polynucleotide sequences encoding an epitope of a polypeptide sequence of the invention (such as, for example, the sequence disclosed in SEQ ID NO:X, or a fragment thereof), polynucleotide sequences of the complementary strand of a polynucleotide sequence encoding an epitope of the invention, and polynucleotide sequences which hybridize to the complementary strand under stringent hybridization conditions or alternatively, under lower stringency hybridization conditions defined *supra*.

deletions. Polynucleotides encoding these polypeptides are also encompassed by the invention.

[0140] Any polypeptide sequence encoded by, for example, the polynucleotide sequences set forth as SEQ ID NO:X or the complement thereof, (presented, for example, in Tables 1 and 2), or the cDNA contained in Clone ID NO:Z, may be analyzed to determine certain preferred regions of the polypeptide. For example, the amino acid sequence of a polypeptide encoded by a polynucleotide sequence of SEQ ID NO:X (e.g., the polypeptide of SEQ ID NO:Y and the polypeptide encoded by the portion of SEQ ID NO:X as defined in columns 8 and 9 of Table 2) or the cDNA contained in Clone ID NO:Z may be analyzed using the default parameters of the DNASTAR computer algorithm (DNASTAR, Inc., 1228 S. Park St., Madison, WI 53715 USA; <http://www.dnastar.com/>).

[0141] Polypeptide regions that may be routinely obtained using the DNASTAR computer algorithm include, but are not limited to, Garnier-Robson alpha-regions, beta-regions, turn-regions, and coil-regions; Chou-Fasman alpha-regions, beta-regions, and turn-regions; Kyte-Doolittle hydrophilic regions and hydrophobic regions; Eisenberg alpha- and beta-amphipathic regions; Karplus-Schulz flexible regions; Emini surface-forming regions; and Jameson-Wolf regions of high antigenic index. Among highly preferred polynucleotides of the invention in this regard are those that encode polypeptides comprising regions that combine several structural features, such as several (e.g., 1, 2, 3 or 4) of the features set out above.

[0142] Additionally, Kyte-Doolittle hydrophilic regions and hydrophobic regions, Emini surface-forming regions, and Jameson-Wolf regions of high antigenic index (i.e., containing four or more contiguous amino acids having an antigenic index of greater than or equal to 1.5, as identified using the default parameters of the Jameson-Wolf program) can routinely be used to determine polypeptide regions that exhibit a high degree of potential for antigenicity. Regions of high antigenicity are determined from data by DNASTAR analysis by choosing values which represent regions of the polypeptide which are likely to be exposed on the surface of the polypeptide in an environment in which antigen recognition may occur in the process of initiation of an immune response.

[0143] Preferred polypeptide fragments of the invention are fragments comprising, or alternatively, consisting of, an amino acid sequence that displays a functional activity (e.g.

NO:X as defined in columns 8 and 9 of Table 2, and/or a polypeptide encoded by the cDNA contained in Clone ID NO:Z). In particular, C-terminal deletions may be described by the general formula 1-n, where n is any whole integer ranging from 6 to q-1, and where n corresponds to the position of amino acid residue in a polypeptide of the invention. Polynucleotides encoding these polypeptides are also encompassed by the invention.

[0137] In addition, any of the above described N- or C-terminal deletions can be combined to produce a N- and C-terminal deleted polypeptide. The invention also provides polypeptides having one or more amino acids deleted from both the amino and the carboxyl termini, which may be described generally as having residues m-n of a polypeptide encoded by SEQ ID NO:X (e.g., including, but not limited to, the preferred polypeptide disclosed as SEQ ID NO:Y and the polypeptide encoded by the portion of SEQ ID NO:X as defined in columns 8 and 9 of Table 2), the cDNA contained in Clone ID NO:Z, and/or the complement thereof, where n and m are integers as described above. Polynucleotides encoding these polypeptides are also encompassed by the invention.

[0138] Also as mentioned above, even if deletion of one or more amino acids from the C-terminus of a protein results in modification or loss of one or more biological functions of the protein, other functional activities (e.g., biological activities, ability to multimerize, ability to bind a ligand) may still be retained. For example the ability of the shortened mutein to induce and/or bind to antibodies which recognize the complete or mature forms of the polypeptide generally will be retained when less than the majority of the residues of the complete or mature polypeptide are removed from the C-terminus. Whether a particular polypeptide lacking C-terminal residues of a complete polypeptide retains such immunologic activities can readily be determined by routine methods described herein and otherwise known in the art. It is not unlikely that a mutein with a large number of deleted C-terminal amino acid residues may retain some biological or immunogenic activities. In fact, peptides composed of as few as six amino acid residues may often evoke an immune response.

[0139] The present application is also directed to proteins containing polypeptides at least 80%, 85%, 90%, 95%, 96%, 97%, 98% or 99% identical to a polypeptide sequence set forth herein. In preferred embodiments, the application is directed to proteins containing polypeptides at least 80%, 85%, 90%, 95%, 96%, 97%, 98% or 99% identical to polypeptides having the amino acid sequence of the specific N- and C-terminal

complete or mature polypeptide are removed from the N-terminus. Whether a particular polypeptide lacking N-terminal residues of a complete polypeptide retains such immunologic activities can readily be determined by routine methods described herein and otherwise known in the art. It is not unlikely that a mutein with a large number of deleted N-terminal amino acid residues may retain some biological or immunogenic activities. In fact, peptides composed of as few as six amino acid residues may often evoke an immune response.

[0134] Accordingly, polypeptide fragments include the secreted protein as well as the mature form. Further preferred polypeptide fragments include the secreted protein or the mature form having a continuous series of deleted residues from the amino or the carboxy terminus, or both. For example, any number of amino acids, ranging from 1-60, can be deleted from the amino terminus of either the secreted polypeptide or the mature form. Similarly, any number of amino acids, ranging from 1-30, can be deleted from the carboxy terminus of the secreted protein or mature form. Furthermore, any combination of the above amino and carboxy terminus deletions are preferred. Similarly, polynucleotides encoding these polypeptide fragments are also preferred.

[0135] The present invention further provides polypeptides having one or more residues deleted from the amino terminus of the amino acid sequence of a polypeptide disclosed herein (e.g., a polypeptide of SEQ ID NO:Y, a polypeptide encoded by the polynucleotide sequence contained in SEQ ID NO:X or the complement thereof, a polypeptide encoded by the portion of SEQ ID NO:X as defined in columns 8 and 9 of Table 2, and/or a polypeptide encoded by the cDNA contained in Clone ID NO:Z). In particular, N-terminal deletions may be described by the general formula m-q, where q is a whole integer representing the total number of amino acid residues in a polypeptide of the invention (e.g., the polypeptide disclosed in SEQ ID NO:Y, or the polypeptide encoded by the portion of SEQ ID NO:X as defined in columns 8 and 9 of Table 2), and m is defined as any integer ranging from 2 to q-6. Polynucleotides encoding these polypeptides are also encompassed by the invention.

[0136] The present invention further provides polypeptides having one or more residues from the carboxy terminus of the amino acid sequence of a polypeptide disclosed herein (e.g., a polypeptide of SEQ ID NO:Y, a polypeptide encoded by the polynucleotide sequence contained in SEQ ID NO:X, a polypeptide encoded by the portion of SEQ ID

alternatively consisting of, from about amino acid number 1-20, 21-40, 41-60, 61-80, 81-100, 101-120, 121-140, 141-160, 161-180, 181-200, 201-220, 221-240, 241-260, 261-280, 281-300, 301-320, 321-340, 341-360, 361-380, 381-400, 401-420, 421-440, 441-460, 461-480, 481-500, 501-520, 521-540, 541-560, 561-580, 581-600, 601-620, 621-640, 641-660, 661-680, 681-700, 701-720, 721-740, 741-760, 761-780, 781-800, 801-820, 821-840, 841-860, 861-880, 881-900, 901-920, 921-940, 941-960, 961-980, 981-1000, 1001-1020, 1021-1040, 1041-1060, 1061-1080, 1081-1100, 1101-1120, 1121-1140, 1141-1160, 1161-1180, 1181-1200, 1201-1220, 1221-1240, 1241-1260, 1261-1280, 1281-1300, 1301-1320, 1321-1340, 1341-1360, 1361-1380, 1381-1400, 1401-1420, 1421-1440, or 1441 to the end of the coding region of cDNA and SEQ ID NO: Y. In a preferred embodiment, polypeptide fragments of the invention include, for example, fragments comprising, or alternatively consisting of, from about amino acid number 1-20, 21-40, 41-60, 61-80, 81-100, 101-120, 121-140, 141-160, 161-180, 181-200, 201-220, 221-240, 241-260, 261-280, 281-300, 301-320, 321-340, 341-360, 361-380, 381-400, 401-420, 421-440, 441-460, 461-480, 481-500, 501-520, 521-540, 541-560, 561-580, 581-600, 601-620, 621-640, 641-660, 661-680, 681-700, 701-720, 721-740, 741-760, 761-780, 781-800, 801-820, 821-840, 841-860, 861-880, 881-900, 901-920, 921-940, 941-960, 961-980, 981-1000, 1001-1020, 1021-1040, 1041-1060, 1061-1080, 1081-1100, 1101-1120, 1121-1140, 1141-1160, 1161-1180, 1181-1200, 1201-1220, 1221-1240, 1241-1260, 1261-1280, 1281-1300, 1301-1320, 1321-1340, 1341-1360, 1361-1380, 1381-1400, 1401-1420, 1421-1440, or 1441 to the end of the coding region of SEQ ID NO:Y. Moreover, polypeptide fragments of the invention may be at least about 10, 15, 20, 25, 30, 35, 40, 45, 50, 55, 60, 65, 70, 75, 80, 85, 90, 100, 110, 120, 130, 140, or 150 amino acids in length. In this context "about" includes the particularly recited ranges or values, or ranges or values larger or smaller by several (5, 4, 3, 2, or 1) amino acids, at either extreme or at both extremes. Polynucleotides encoding these polypeptide fragments are also encompassed by the invention.

[0133] Even if deletion of one or more amino acids from the N-terminus of a protein results in modification or loss of one or more biological functions of the protein, other functional activities (e.g., biological activities, ability to multimerize, ability to bind a ligand) may still be retained. For example, the ability of shortened muteins to induce and/or bind to antibodies which recognize the complete or mature forms of the polypeptides generally will be retained when less than the majority of the residues of the

2501-2550, 2551-2600, 2601-2650, 2651-2700, 2701-2750, 2751-2800, 2801-2850, 2851-2900, 2901-2950, 2951-3000, 3001-3050, 3051-3100, 3101-3150, 3151-3200, 3201-3250, 3251-3300, 3301-3350, 3351-3400, 3401-3450, 3451-3500, 3501-3550, 3551-3600, 3601-3650, 3651-3700, 3701-3750, 3751-3800, 3801-3850, 3851-3900, 3901-3950, 3951-4000, 4001-4050, 4051-4100, 4101-4150, 4151-4200, 4201-4250, 4251-4300, 4301-4350, 4351-4400, 4401-4450, 4451-4500, 4501-4550, 4551-4600, 4601-4650, 4651-4700, 4701-4750, 4751-4800, 4801-4850, 4851-4900, 4901-4950, 4951-5000, 5001-5050, 5051-5100, 5101-5150, 5151-5200, 5201-5250, 5251-5300, 5301-5350, 5351-5400, 5401-5450, 5451-5500, 5501-5550, 5551-5600, 5601-5650, 5651-5700, 5701-5750, 5751-5800, 5801-5850, 5851-5900, 5901-5950, 5951-6000, 6001-6050, 6051-6100, 6101-6150, 6151-6200, 6201-6250, 6251-6300, 6301-6350, 6351-6400, 6401-6450, 6451-6500, 6501-6550, 6551-6600, 6601-6650, 6651-6700, 6701-6750, 6751-6800, 6801-6850, 6851-6900, 6901-6950, 6951-7000, 7001-7050, 7051-7100, 7101-7150, 7151-7200, 7201-7250, 7251-7300 or 7301 to the end of the cDNA sequence contained in Clone ID NO:Z, or the complementary strand thereto.

In this context "about" includes the particularly recited range or a range larger or smaller by several (5, 4, 3, 2, or 1) nucleotides, at either terminus or at both termini. Preferably, these fragments encode a polypeptide which has a functional activity (e.g., biological activity). More preferably, these polynucleotides can be used as probes or primers as discussed herein. Polynucleotides which hybridize to one or more of these polynucleotides under stringent hybridization conditions or alternatively, under lower stringency conditions are also encompassed by the invention, as are polypeptides encoded by these polynucleotides.

[0132] In the present invention, a "polypeptide fragment" refers to an amino acid sequence which is a portion of that contained in SEQ ID NO:Y, a portion of an amino acid sequence encoded by the portion of SEQ ID NO:X as defined in columns 8 and 9 of Table 2, a portion of an amino acid sequence encoded by the polynucleotide sequence of SEQ ID NO:X, a portion of an amino acid sequence encoded by the complement of the polynucleotide sequence in SEQ ID NO:X, and/or a portion of an amino acid sequence encoded by the cDNA contained in Clone ID NO:Z. Protein (polypeptide) fragments may be "free-standing," or comprised within a larger polypeptide of which the fragment forms a part or region, most preferably as a single continuous region. Representative examples of polypeptide fragments of the invention, include, for example, fragments comprising, or

1351-1400, 1401-1450, 1451-1500, 1501-1550, 1551-1600, 1601-1650, 1651-1700, 1701-1750, 1751-1800, 1801-1850, 1851-1900, 1901-1950, 1951-2000, 2001-2050, 2051-2100, 2101-2150, 2151-2200, 2201-2250, 2251-2300, 2301-2350, 2351-2400, 2401-2450, 2451-2500, 2501-2550, 2551-2600, 2601-2650, 2651-2700, 2701-2750, 2751-2800, 2801-2850, 2851-2900, 2901-2950, 2951-3000, 3001-3050, 3051-3100, 3101-3150, 3151-3200, 3201-3250, 3251-3300, 3301-3350, 3351-3400, 3401-3450, 3451-3500, 3501-3550, 3551-3600, 3601-3650, 3651-3700, 3701-3750, 3751-3800, 3801-3850, 3851-3900, 3901-3950, 3951-4000, 4001-4050, 4051-4100, 4101-4150, 4151-4200, 4201-4250, 4251-4300, 4301-4350, 4351-4400, 4401-4450, 4451-4500, 4501-4550, 4551-4600, 4601-4650, 4651-4700, 4701-4750, 4751-4800, 4801-4850, 4851-4900, 4901-4950, 4951-5000, 5001-5050, 5051-5100, 5101-5150, 5151-5200, 5201-5250, 5251-5300, 5301-5350, 5351-5400, 5401-5450, 5451-5500, 5501-5550, 5551-5600, 5601-5650, 5651-5700, 5701-5750, 5751-5800, 5801-5850, 5851-5900, 5901-5950, 5951-6000, 6001-6050, 6051-6100, 6101-6150, 6151-6200, 6201-6250, 6251-6300, 6301-6350, 6351-6400, 6401-6450, 6451-6500, 6501-6550, 6551-6600, 6601-6650, 6651-6700, 6701-6750, 6751-6800, 6801-6850, 6851-6900, 6901-6950, 6951-7000, 7001-7050, 7051-7100, 7101-7150, 7151-7200, 7201-7250, 7251-7300 or 7301 to the end of SEQ ID NO:X, or the complementary strand thereto. In this context "about" includes the particularly recited range or a range larger or smaller by several (5, 4, 3, 2, or 1) nucleotides, at either terminus or at both termini. Preferably, these fragments encode a polypeptide which has a functional activity (e.g., biological activity). More preferably, these polynucleotides can be used as probes or primers as discussed herein. Polynucleotides which hybridize to one or more of these polynucleotides under stringent hybridization conditions or alternatively, under lower stringency conditions are also encompassed by the invention, as are polypeptides encoded by these polynucleotides.

[0131] Further representative examples of polynucleotide fragments of the invention comprise, or alternatively consist of, a sequence from about nucleotide number 1-50, 51-100, 101-150, 151-200, 201-250, 251-300, 301-350, 351-400, 401-450, 451-500, 501-550, 551-600, 601-650, 651-700, 701-750, 751-800, 801-850, 851-900, 901-950, 951-1000, 1001-1050, 1051-1100, 1101-1150, 1151-1200, 1201-1250, 1251-1300, 1301-1350, 1351-1400, 1401-1450, 1451-1500, 1501-1550, 1551-1600, 1601-1650, 1651-1700, 1701-1750, 1751-1800, 1801-1850, 1851-1900, 1901-1950, 1951-2000, 2001-2050, 2051-2100, 2101-2150, 2151-2200, 2201-2250, 2251-2300, 2301-2350, 2351-2400, 2401-2450, 2451-2500,

[0128] The present invention is also directed to polynucleotide fragments of the polynucleotides (nucleic acids) of the invention. In the present invention, a "polynucleotide fragment" refers to a polynucleotide having a nucleic acid sequence which, for example: is a portion of the cDNA contained in Clone ID NO:Z or the complementary strand thereto; is a portion of the polynucleotide sequence encoding the polypeptide encoded by the cDNA contained in Clone ID NO:Z or the complementary strand thereto; is a portion of a polynucleotide sequence encoding the amino acid sequence encoded by the region of SEQ ID NO:X as defined in columns 8 and 9 of Table 2 or the complementary strand thereto; is a portion of the polynucleotide sequence of SEQ ID NO:X as defined in columns 8 and 9 of Table 2 or the complementary strand thereto; is a portion of the polynucleotide sequence in SEQ ID NO:X or the complementary strand thereto; is a polynucleotide sequence encoding a portion of the polypeptide of SEQ ID NO:Y; is a polynucleotide sequence encoding a portion of a polypeptide encoded by SEQ ID NO:X; or is a polynucleotide sequence encoding a portion of a polypeptide encoded by the complement of the polynucleotide sequence in SEQ ID NO:X.

[0129] The polynucleotide fragments of the invention are preferably at least about 15 nt, and more preferably at least about 20 nt, still more preferably at least about 30 nt, and even more preferably, at least about 40 nt, at least about 50 nt, at least about 75 nt, or at least about 150 nt in length. A fragment "at least 20 nt in length," for example, is intended to include 20 or more contiguous bases from the cDNA sequence contained in Clone ID NO:Z, or the nucleotide sequence shown in SEQ ID NO:X or the complementary strand thereto. In this context "about" includes the particularly recited value or a value larger or smaller by several (5, 4, 3, 2, or 1) nucleotides, at either terminus or at both termini. These nucleotide fragments have uses that include, but are not limited to, as diagnostic probes and primers as discussed herein. Of course, larger fragments (e.g., at least 160, 170, 180, 190, 200, 250, 500, 600, 1000, or 2000 nucleotides in length) are also encompassed by the invention.

[0130] Moreover, representative examples of polynucleotide fragments of the invention comprise, or alternatively consist of, a sequence from about nucleotide number 1-50, 51-100, 101-150, 151-200, 201-250, 251-300, 301-350, 351-400, 401-450, 451-500, 501-550, 551-600, 601-650, 651-700, 701-750, 751-800, 801-850, 851-900, 901-950, 951-1000, 1001-1050, 1051-1100, 1101-1150, 1151-1200, 1201-1250, 1251-1300, 1301-1350,

formulations both reduces activity and increases clearance due to the aggregate's immunogenic activity. See Pinckard et al., Clin. Exp. Immunol. 2:331-340 (1967); Robbins et al., Diabetes 36: 838-845 (1987); Cleland et al., Crit. Rev. Therapeutic Drug Carrier Systems 10:307-377 (1993).

[0126] A further embodiment of the invention relates to polypeptides which comprise the amino acid sequence of a polypeptide having an amino acid sequence which contains at least one amino acid substitution, but not more than 50 amino acid substitutions, even more preferably, not more than 40 amino acid substitutions, still more preferably, not more than 30 amino acid substitutions, and still even more preferably, not more than 20 amino acid substitutions from a polypeptide sequence disclosed herein. Of course it is highly preferable for a polypeptide to have an amino acid sequence which comprises the amino acid sequence of a polypeptide of SEQ ID NO:Y, an amino acid sequence encoded by SEQ ID NO:X, an amino acid sequence encoded by the portion of SEQ ID NO:X as defined in columns 8 and 9 of Table 2, an amino acid sequence encoded by the complement of SEQ ID NO:X, and/or an amino acid sequence encoded by cDNA contained in Clone ID NO:Z which contains, in order of ever-increasing preference, at least one, but not more than 10, 9, 8, 7, 6, 5, 4, 3, 2 or 1 amino acid substitutions.

[0127] In specific embodiments, the polypeptides of the invention comprise, or alternatively, consist of, fragments or variants of a reference amino acid sequence selected from: (a) the amino acid sequence of SEQ ID NO:Y or fragments thereof (e.g., the mature form and/or other fragments described herein); (b) the amino acid sequence encoded by SEQ ID NO:X or fragments thereof; (c) the amino acid sequence encoded by the complement of SEQ ID NO:X or fragments thereof; (d) the amino acid sequence encoded by the portion of SEQ ID NO:X as defined in columns 8 and 9 of Table 2 or fragments thereof; and (e) the amino acid sequence encoded by cDNA contained in Clone ID NO:Z or fragments thereof; wherein the fragments or variants have 1-5, 5-10, 5-25, 5-50, 10-50 or 50-150, amino acid residue additions, substitutions, and/or deletions when compared to the reference amino acid sequence. In preferred embodiments, the amino acid substitutions are conservative. Polynucleotides encoding these polypeptides are also encompassed by the invention.

Polynucleotide and Polypeptide Fragments

example, site directed mutagenesis or alanine-scanning mutagenesis (introduction of single alanine mutations at every residue in the molecule) can be used. See Cunningham and Wells, Science 244:1081-1085 (1989). The resulting mutant molecules can then be tested for biological activity.

[0124] As the authors state, these two strategies have revealed that proteins are surprisingly tolerant of amino acid substitutions. The authors further indicate which amino acid changes are likely to be permissive at certain amino acid positions in the protein. For example, most buried (within the tertiary structure of the protein) amino acid residues require nonpolar side chains, whereas few features of surface side chains are generally conserved. Moreover, tolerated conservative amino acid substitutions involve replacement of the aliphatic or hydrophobic amino acids Ala, Val, Leu and Ile; replacement of the hydroxyl residues Ser and Thr; replacement of the acidic residues Asp and Glu; replacement of the amide residues Asn and Gln, replacement of the basic residues Lys, Arg, and His; replacement of the aromatic residues Phe, Tyr, and Trp, and replacement of the small-sized amino acids Ala, Ser, Thr, Met, and Gly. Besides conservative amino acid substitution, variants of the present invention include (i) substitutions with one or more of the non-conserved amino acid residues, where the substituted amino acid residues may or may not be one encoded by the genetic code, or (ii) substitutions with one or more of the amino acid residues having a substituent group, or (iii) fusion of the mature polypeptide with another compound, such as a compound to increase the stability and/or solubility of the polypeptide (for example, polyethylene glycol), (iv) fusion of the polypeptide with additional amino acids, such as, for example, an IgG Fc fusion region peptide, serum albumin (preferably human serum albumin) or a fragment thereof, or leader or secretory sequence, or a sequence facilitating purification, or (v) fusion of the polypeptide with another compound, such as albumin (including but not limited to recombinant albumin (see, e.g., U.S. Patent No. 5,876,969, issued March 2, 1999, EP Patent 0 413 622, and U.S. Patent No. 5,766,883, issued June 16, 1998, herein incorporated by reference in their entirety)). Such variant polypeptides are deemed to be within the scope of those skilled in the art from the teachings herein.

[0125] For example, polypeptide variants containing amino acid substitutions of charged amino acids with other charged or neutral amino acids may produce proteins with improved characteristics, such as less aggregation. Aggregation of pharmaceutical

and fragments, variants and derivatives thereof to elicit polypeptide related biological activity (either *in vitro* or *in vivo*). Other methods will be known to the skilled artisan and are within the scope of the invention.

[0120] Of course, due to the degeneracy of the genetic code, one of ordinary skill in the art will immediately recognize that a large number of the nucleic acid molecules having a sequence at least 80%, 85%, 90%, 95%, 96%, 97%, 98%, 99%, or 100% identical to, for example, the nucleic acid sequence of the cDNA contained in Clone ID NO:Z, the nucleic acid sequence referred to in Table 1 (SEQ ID NO:X), the nucleic acid sequence disclosed in Table 2 (e.g., the nucleic acid sequence delineated in columns 8 and 9) or fragments thereof, will encode polypeptides "having functional activity." In fact, since degenerate variants of any of these nucleotide sequences all encode the same polypeptide, in many instances, this will be clear to the skilled artisan even without performing the above described comparison assay. It will be further recognized in the art that, for such nucleic acid molecules that are not degenerate variants, a reasonable number will also encode a polypeptide having functional activity. This is because the skilled artisan is fully aware of amino acid substitutions that are either less likely or not likely to significantly effect protein function (e.g., replacing one aliphatic amino acid with a second aliphatic amino acid), as further described below.

[0121] For example, guidance concerning how to make phenotypically silent amino acid substitutions is provided in Bowie et al., "Deciphering the Message in Protein Sequences: Tolerance to Amino Acid Substitutions," *Science* 247:1306-1310 (1990), wherein the authors indicate that there are two main strategies for studying the tolerance of an amino acid sequence to change.

[0122] The first strategy exploits the tolerance of amino acid substitutions by natural selection during the process of evolution. By comparing amino acid sequences in different species, conserved amino acids can be identified. These conserved amino acids are likely important for protein function. In contrast, the amino acid positions where substitutions have been tolerated by natural selection indicates that these positions are not critical for protein function. Thus, positions tolerating amino acid substitution could be modified while still maintaining biological activity of the protein.

[0123] The second strategy uses genetic engineering to introduce amino acid changes at specific positions of a cloned gene to identify regions critical for protein function. For

antigenicity [ability to bind (or compete with a polypeptide of the invention for binding) to an anti-polypeptide of the invention antibody], immunogenicity (ability to generate antibody which binds to a specific polypeptide of the invention), ability to form multimers with polypeptides of the invention, and ability to bind to a receptor or ligand for a polypeptide of the invention.

[0116] The functional activity of the polypeptides, and fragments, variants and derivatives of the invention, can be assayed by various methods.

[0117] For example, in one embodiment where one is assaying for the ability to bind or compete with a full-length polypeptide of the present invention for binding to an anti-polypeptide antibody, various immunoassays known in the art can be used, including but not limited to, competitive and non-competitive assay systems using techniques such as radioimmunoassays, ELISA (enzyme linked immunosorbent assay), "sandwich" immunoassays, immunoradiometric assays, gel diffusion precipitation reactions, immunodiffusion assays, in situ immunoassays (using colloidal gold, enzyme or radioisotope labels, for example), western blots, precipitation reactions, agglutination assays (e.g., gel agglutination assays, hemagglutination assays), complement fixation assays, immunofluorescence assays, protein A assays, and immunoelectrophoresis assays, etc. In one embodiment, antibody binding is detected by detecting a label on the primary antibody. In another embodiment, the primary antibody is detected by detecting binding of a secondary antibody or reagent to the primary antibody. In a further embodiment, the secondary antibody is labeled. Many means are known in the art for detecting binding in an immunoassay and are within the scope of the present invention.

[0118] In another embodiment, where a ligand is identified, or the ability of a polypeptide fragment, variant or derivative of the invention to multimerize is being evaluated, binding can be assayed, e.g., by means well-known in the art, such as, for example, reducing and non-reducing gel chromatography, protein affinity chromatography, and affinity blotting. See generally, Phizicky et al., Microbiol. Rev. 59:94-123 (1995). In another embodiment, the ability of physiological correlates of a polypeptide of the present invention to bind to a substrate(s) of the polypeptide of the invention can be routinely assayed using techniques known in the art.

[0119] In addition, assays described herein (see Examples) and otherwise known in the art may routinely be applied to measure the ability of polypeptides of the present invention

functions, other biological activities may still be retained. For example, the ability of a deletion variant to induce and/or to bind antibodies which recognize the secreted form will likely be retained when less than the majority of the residues of the secreted form are removed from the N-terminus or C-terminus. Whether a particular polypeptide lacking N- or C-terminal residues of a protein retains such immunogenic activities can readily be determined by routine methods described herein and otherwise known in the art.

[0113] Thus, the invention further includes polypeptide variants which show a functional activity (e.g., biological activity) of the polypeptides of the invention. Such variants include deletions, insertions, inversions, repeats, and substitutions selected according to general rules known in the art so as have little effect on activity.

[0114] The present application is directed to nucleic acid molecules at least 80%, 85%, 90%, 95%, 96%, 97%, 98%, 99% or 100% identical to the nucleic acid sequences disclosed herein, (e.g., encoding a polypeptide having the amino acid sequence of an N and/or C terminal deletion), irrespective of whether they encode a polypeptide having functional activity. This is because even where a particular nucleic acid molecule does not encode a polypeptide having functional activity, one of skill in the art would still know how to use the nucleic acid molecule, for instance, as a hybridization probe or a polymerase chain reaction (PCR) primer. Uses of the nucleic acid molecules of the present invention that do not encode a polypeptide having functional activity include, inter alia, (1) isolating a gene or allelic or splice variants thereof in a cDNA library; (2) *in situ* hybridization (e.g., "FISH") to metaphase chromosomal spreads to provide precise chromosomal location of the gene, as described in Verma et al., *Human Chromosomes: A Manual of Basic Techniques*, Pergamon Press, New York (1988); (3) Northern Blot analysis for detecting mRNA expression in specific tissues (e.g., normal or diseased tissues); and (4) *in situ* hybridization (e.g., histochemistry) for detecting mRNA expression in specific tissues (e.g., normal or diseased tissues).

[0115] Preferred, however, are nucleic acid molecules having sequences at least 80%, 85%, 90%, 95%, 96%, 97%, 98%, 99% or 100% identical to the nucleic acid sequences disclosed herein, which do, in fact, encode a polypeptide having functional activity. By a polypeptide having "functional activity" is meant, a polypeptide capable of displaying one or more known functional activities associated with a full-length (complete) protein of the invention. Such functional activities include, but are not limited to, biological activity,

30, less than 20, less than 10, or 5-50, 5-25, 5-10, 1-5, or 1-2 amino acids are substituted, deleted, or added in any combination are also preferred. Polynucleotide variants can be produced for a variety of reasons, e.g., to optimize codon expression for a particular host (change codons in the human mRNA to those preferred by a bacterial host such as *E. coli*).

[0109] Naturally occurring variants are called "allelic variants," and refer to one of several alternate forms of a gene occupying a given locus on a chromosome of an organism. (Genes II, Lewin, B., ed., John Wiley & Sons, New York (1985)). These allelic variants can vary at either the polynucleotide and/or polypeptide level and are included in the present invention. Alternatively, non-naturally occurring variants may be produced by mutagenesis techniques or by direct synthesis.

[0110] Using known methods of protein engineering and recombinant DNA technology, variants may be generated to improve or alter the characteristics of the polypeptides of the present invention. For instance, one or more amino acids can be deleted from the N-terminus or C-terminus of the polypeptide of the present invention without substantial loss of biological function. As an example, Ron et al. (*J. Biol. Chem.* 268: 2984-2988 (1993)) reported variant KGF proteins having heparin binding activity even after deleting 3, 8, or 27 amino-terminal amino acid residues. Similarly, Interferon-gamma exhibited up to ten times higher activity after deleting 8-10 amino acid residues from the carboxy terminus of this protein. (Dobeli et al., *J. Biotechnology* 7:199-216 (1988).)

[0111] Moreover, ample evidence demonstrates that variants often retain a biological activity similar to that of the naturally occurring protein. For example, Gayle and coworkers (*J. Biol. Chem.* 268:22105-22111 (1993)) conducted extensive mutational analysis of human cytokine IL-1a. They used random mutagenesis to generate over 3,500 individual IL-1a mutants that averaged 2.5 amino acid changes per variant over the entire length of the molecule. Multiple mutations were examined at every possible amino acid position. The investigators found that "[m]ost of the molecule could be altered with little effect on either [binding or biological activity]." In fact, only 23 unique amino acid sequences, out of more than 3,500 nucleotide sequences examined, produced a protein that significantly differed in activity from wild-type.

[0112] Furthermore, even if deleting one or more amino acids from the N-terminus or C-terminus of a polypeptide results in modification or loss of one or more biological

N- and C-terminal of the subject sequence, which are not matched/aligned with a corresponding subject residue, as a percent of the total bases of the query sequence. Whether a residue is matched/aligned is determined by results of the FASTDB sequence alignment. This percentage is then subtracted from the percent identity, calculated by the above FASTDB program using the specified parameters, to arrive at a final percent identity score. This final percent identity score is what is used for the purposes of the present invention. Only residues to the N- and C-termini of the subject sequence, which are not matched/aligned with the query sequence, are considered for the purposes of manually adjusting the percent identity score. That is, only query residue positions outside the farthest N- and C- terminal residues of the subject sequence.

[0107] For example, a 90 amino acid residue subject sequence is aligned with a 100 residue query sequence to determine percent identity. The deletion occurs at the N-terminus of the subject sequence and therefore, the FASTDB alignment does not show a matching/alignment of the first 10 residues at the N-terminus. The 10 unpaired residues represent 10% of the sequence (number of residues at the N- and C- termini not matched/total number of residues in the query sequence) so 10% is subtracted from the percent identity score calculated by the FASTDB program. If the remaining 90 residues were perfectly matched the final percent identity would be 90%. In another example, a 90 residue subject sequence is compared with a 100 residue query sequence. This time the deletions are internal deletions so there are no residues at the N- or C-termini of the subject sequence which are not matched/aligned with the query. In this case the percent identity calculated by FASTDB is not manually corrected. Once again, only residue positions outside the N- and C-terminal ends of the subject sequence, as displayed in the FASTDB alignment, which are not matched/aligned with the query sequence are manually corrected for. No other manual corrections are to be made for the purposes of the present invention.

[0108] The polynucleotide variants of the invention may contain alterations in the coding regions, non-coding regions, or both. Especially preferred are polynucleotide variants containing alterations which produce silent substitutions, additions, or deletions, but do not alter the properties or activities of the encoded polypeptide. Nucleotide variants produced by silent substitutions due to the degeneracy of the genetic code are preferred. Moreover, polypeptide variants in which less than 50, less than 40, less than

polypeptide having an amino acid sequence at least 95% identical to a query amino acid sequence, up to 5% of the amino acid residues in the subject sequence may be inserted, deleted, (indels) or substituted with another amino acid. These alterations of the reference sequence may occur at the amino or carboxy terminal positions of the reference amino acid sequence or anywhere between those terminal positions, interspersed either individually among residues in the reference sequence or in one or more contiguous groups within the reference sequence.

[0105] As a practical matter, whether any particular polypeptide is at least 80%, 85%, 90%, 95%, 96%, 97%, 98% or 99% identical to, for instance, the amino acid sequence of a polypeptide referred to in Table 1 (e.g., the amino acid sequence identified in column 6) or Table 2 (e.g., the amino acid sequence of the polypeptide encoded by the polynucleotide sequence defined in columns 8 and 9 of Table 2) or a fragment thereof, the amino acid sequence of the polypeptide encoded by the nucleotide sequence in SEQ ID NO:X or a fragment thereof, or the amino acid sequence of the polypeptide encoded by cDNA contained in Clone ID NO:Z, or a fragment thereof, can be determined conventionally using known computer programs. A preferred method for determining the best overall match between a query sequence (a sequence of the present invention) and a subject sequence, also referred to as a global sequence alignment, can be determined using the FASTDB computer program based on the algorithm of Brutlag et al. (Comp. App. Biosci.6:237-245 (1990)). In a sequence alignment the query and subject sequences are either both nucleotide sequences or both amino acid sequences. The result of said global sequence alignment is expressed as percent identity. Preferred parameters used in a FASTDB amino acid alignment are: Matrix=PAM 0, k-tuple=2, Mismatch Penalty=1, Joining Penalty=20, Randomization Group Length=0, Cutoff Score=1, Window Size=sequence length, Gap Penalty=5, Gap Size Penalty=0.05, Window Size=500 or the length of the subject amino acid sequence, whichever is shorter.

[0106] If the subject sequence is shorter than the query sequence due to N- or C-terminal deletions, not because of internal deletions, a manual correction must be made to the results. This is because the FASTDB program does not account for N- and C-terminal truncations of the subject sequence when calculating global percent identity. For subject sequences truncated at the N- and C-termini, relative to the query sequence, the percent identity is corrected by calculating the number of residues of the query sequence that are

results. This is because the FASTDB program does not account for 5' and 3' truncations of the subject sequence when calculating percent identity. For subject sequences truncated at the 5' or 3' ends, relative to the query sequence, the percent identity is corrected by calculating the number of bases of the query sequence that are 5' and 3' of the subject sequence, which are not matched/aligned, as a percent of the total bases of the query sequence. Whether a nucleotide is matched/aligned is determined by results of the FASTDB sequence alignment. This percentage is then subtracted from the percent identity, calculated by the above FASTDB program using the specified parameters, to arrive at a final percent identity score. This corrected score is what is used for the purposes of the present invention. Only bases outside the 5' and 3' bases of the subject sequence, as displayed by the FASTDB alignment, which are not matched/aligned with the query sequence, are calculated for the purposes of manually adjusting the percent identity score.

[0103] For example, a 90 base subject sequence is aligned to a 100 base query sequence to determine percent identity. The deletions occur at the 5' end of the subject sequence and therefore, the FASTDB alignment does not show a matched/alignment of the first 10 bases at 5' end. The 10 unpaired bases represent 10% of the sequence (number of bases at the 5' and 3' ends not matched/total number of bases in the query sequence) so 10% is subtracted from the percent identity score calculated by the FASTDB program. If the remaining 90 bases were perfectly matched the final percent identity would be 90%. In another example, a 90 base subject sequence is compared with a 100 base query sequence. This time the deletions are internal deletions so that there are no bases on the 5' or 3' of the subject sequence which are not matched/aligned with the query. In this case the percent identity calculated by FASTDB is not manually corrected. Once again, only bases 5' and 3' of the subject sequence which are not matched/aligned with the query sequence are manually corrected for. No other manual corrections are to be made for the purposes of the present invention.

[0104] By a polypeptide having an amino acid sequence at least, for example, 95% "identical" to a query amino acid sequence of the present invention, it is intended that the amino acid sequence of the subject polypeptide is identical to the query sequence except that the subject polypeptide sequence may include up to five amino acid alterations per each 100 amino acids of the query amino acid sequence. In other words, to obtain a

fragments described herein). Further proteins encoded by polynucleotides which hybridize to the complement of the nucleic acid molecules encoding these amino acid sequences under stringent hybridization conditions or alternatively, under lower stringency conditions, are also encompassed by the invention, as are the polynucleotides encoding these proteins.

[0100] By a nucleic acid having a nucleotide sequence at least, for example, 95% "identical" to a reference nucleotide sequence of the present invention, it is intended that the nucleotide sequence of the nucleic acid is identical to the reference sequence except that the nucleotide sequence may include up to five point mutations per each 100 nucleotides of the reference nucleotide sequence encoding the polypeptide. In other words, to obtain a nucleic acid having a nucleotide sequence at least 95% identical to a reference nucleotide sequence, up to 5% of the nucleotides in the reference sequence may be deleted or substituted with another nucleotide, or a number of nucleotides up to 5% of the total nucleotides in the reference sequence may be inserted into the reference sequence. The query sequence may be an entire sequence referred to in Table 1 or 2 as the ORF (open reading frame), or any fragment specified as described herein.

[0101] As a practical matter, whether any particular nucleic acid molecule or polypeptide is at least 80%, 85%, 90%, 95%, 96%, 97%, 98% or 99% identical to a nucleotide sequence of the present invention can be determined conventionally using known computer programs. A preferred method for determining the best overall match between a query sequence (a sequence of the present invention) and a subject sequence, also referred to as a global sequence alignment, can be determined using the FASTDB computer program based on the algorithm of Brutlag et al. (Comp. App. Biosci. 6:237-245 (1990)). In a sequence alignment the query and subject sequences are both DNA sequences. An RNA sequence can be compared by converting U's to T's. The result of said global sequence alignment is expressed as percent identity. Preferred parameters used in a FASTDB alignment of DNA sequences to calculate percent identity are: Matrix=Unitary, k-tuple=4, Mismatch Penalty=1, Joining Penalty=30, Randomization Group Length=0, Cutoff Score=1, Gap Penalty=5, Gap Size Penalty 0.05, Window Size=500 or the length of the subject nucleotide sequence, whichever is shorter.

[0102] If the subject sequence is shorter than the query sequence because of 5' or 3' deletions, not because of internal deletions, a manual correction must be made to the

acid molecules under stringent hybridization conditions or alternatively, under lower stringency conditions, are also encompassed by the invention, as are polypeptides encoded by these polynucleotides and nucleic acids.

[0097] In a preferred embodiment, the invention encompasses nucleic acid molecules which comprise, or alternatively, consist of a polynucleotide which hybridizes under stringent hybridization conditions, or alternatively, under lower stringency conditions, to a polynucleotide in (a), (b), (c), (d), (e), (f), (g), (h), or (i), above, as are polypeptides encoded by these polynucleotides. In another preferred embodiment, polynucleotides which hybridize to the complement of these nucleic acid molecules under stringent hybridization conditions, or alternatively, under lower stringency conditions, are also encompassed by the invention, as are polypeptides encoded by these polynucleotides.

[0098] In another embodiment, the invention provides a purified protein comprising, or alternatively consisting of, a polypeptide having an amino acid sequence selected from the group consisting of: (a) the complete amino acid sequence of SEQ ID NO:Y or the complete amino acid sequence encoded by the cDNA in Clone ID NO:Z; (b) the amino acid sequence of a mature form of a polypeptide having the amino acid sequence of SEQ ID NO:Y or the amino acid sequence encoded by the cDNA in Clone ID NO:Z; (c) the amino acid sequence of a biologically active fragment of a polypeptide having the complete amino acid sequence of SEQ ID NO:Y or the complete amino acid sequence encoded by the cDNA in Clone ID NO:Z; and (d) the amino acid sequence of an antigenic fragment of a polypeptide having the complete amino acid sequence of SEQ ID NO:Y or the complete amino acid sequence encoded by the cDNA in Clone ID NO:Z.

[0099] The present invention is also directed to proteins which comprise, or alternatively consist of, an amino acid sequence which is at least 80%, 85%, 90%, 95%, 96%, 97%, 98%, 99% or 100%, identical to, for example, any of the amino acid sequences in (a), (b), (c), or (d), above, the amino acid sequence shown in SEQ ID NO:Y, the amino acid sequence encoded by the cDNA contained in Clone ID NO:Z, the amino acid sequence of the polypeptide encoded by the nucleotide sequence in SEQ ID NO:X as defined in columns 8 and 9 of Table 2, the amino acid sequence as defined in column 7 of Table 1, an amino acid sequence encoded by the nucleotide sequence in SEQ ID NO:X, and an amino acid sequence encoded by the complement of the polynucleotide sequence in SEQ ID NO:X. Fragments of these polypeptides are also provided (e.g., those

polypeptide; (e) a nucleotide sequence in SEQ ID NO:X or the cDNA sequence of Clone ID NO:Z, which encodes an antigenic fragment of a polypeptide; (f) a nucleotide sequence encoding a polypeptide comprising the complete amino acid sequence of SEQ ID NO:Y or the complete amino acid sequence encoded by the cDNA in Clone ID NO:Z; (g) a nucleotide sequence encoding a mature polypeptide of the amino acid sequence of SEQ ID NO:Y or the amino acid sequence encoded by the cDNA in Clone ID NO:Z; (h) a nucleotide sequence encoding a biologically active fragment of a polypeptide having the complete amino acid sequence of SEQ ID NO:Y or the complete amino acid sequence encoded by the cDNA in Clone ID NO:Z; (i) a nucleotide sequence encoding an antigenic fragment of a polypeptide having the complete amino acid sequence of SEQ ID NO:Y or the complete amino acid sequence encoded by the cDNA in Clone ID NO:Z; and (j) a nucleotide sequence complementary to any of the nucleotide sequences in (a), (b), (c), (d), (e), (f), (g), (h), or (i) above.

[0096] The present invention is also directed to nucleic acid molecules which comprise, or alternatively consist of, a nucleotide sequence which is at least 80%, 85%, 90%, 95%, 96%, 97%, 98%, 99% or 100%, identical to, for example, any of the nucleotide sequences in (a), (b), (c), (d), (e), (f), (g), (h), (i), or (j) above, the nucleotide coding sequence in SEQ ID NO:X or the complementary strand thereto, the nucleotide coding sequence of the cDNA contained in Clone ID NO:Z or the complementary strand thereto, a nucleotide sequence encoding the polypeptide of SEQ ID NO:Y, a nucleotide sequence encoding a polypeptide sequence encoded by the nucleotide sequence in SEQ ID NO:X, a polypeptide sequence encoded by the complement of the polynucleotide sequence in SEQ ID NO:X, a nucleotide sequence encoding the polypeptide encoded by the cDNA contained in Clone ID NO:Z, the nucleotide coding sequence in SEQ ID NO:X as defined in columns 8 and 9 of Table 2 or the complementary strand thereto, a nucleotide sequence encoding the polypeptide encoded by the nucleotide sequence in SEQ ID NO:X as defined in columns 8 and 9 of Table 2 or the complementary strand thereto, the nucleotide sequence in SEQ ID NO:X encoding the polypeptide sequence as defined in column 7 of Table 1 or the complementary strand thereto, nucleotide sequences encoding the polypeptide as defined in column 7 of Table 1 or the complementary strand thereto, and/or polynucleotide fragments of any of these nucleic acid molecules (e.g., those fragments described herein). Polynucleotides which hybridize to the complement of these nucleic

Polynucleotide and Polypeptide Variants

[0092] The present invention is directed to variants of the polynucleotide sequence disclosed in SEQ ID NO:X or the complementary strand thereto, nucleotide sequences encoding the polypeptide of SEQ ID NO:Y, the nucleotide sequence of SEQ ID NO:X encoding the polypeptide sequence as defined in column 7 of Table 1, nucleotide sequences encoding the polypeptide as defined in column 7 of Table 1, the nucleotide sequence as defined in columns 8 and 9 of Table 2, nucleotide sequences encoding the polypeptide encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2, the cDNA sequence contained in Clone ID NO:Z, and/or nucleotide sequences encoding the polypeptide encoded by the cDNA sequence contained in Clone ID NO:Z.

[0093] The present invention also encompasses variants of the polypeptide sequence disclosed in SEQ ID NO:Y, the polypeptide sequence as defined in column 7 of Table 1, a polypeptide sequence encoded by the polynucleotide sequence in SEQ ID NO:X, a polypeptide sequence encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2, a polypeptide sequence encoded by the complement of the polynucleotide sequence in SEQ ID NO:X, and/or a polypeptide sequence encoded by the cDNA sequence contained in Clone ID NO:Z.

[0094] "Variant" refers to a polynucleotide or polypeptide differing from the polynucleotide or polypeptide of the present invention, but retaining essential properties thereof. Generally, variants are overall closely similar, and, in many regions, identical to the polynucleotide or polypeptide of the present invention.

[0095] Thus, one aspect of the invention provides an isolated nucleic acid molecule comprising, or alternatively consisting of, a polynucleotide having a nucleotide sequence selected from the group consisting of: (a) a nucleotide sequence described in SEQ ID NO:X or contained in the cDNA sequence of Clone ID NO:Z; (b) a nucleotide sequence in SEQ ID NO:X or the cDNA in Clone ID NO:Z which encodes the complete amino acid sequence of SEQ ID NO:Y or the complete amino acid sequence encoded by the cDNA in Clone ID NO:Z; (c) a nucleotide sequence in SEQ ID NO:X or the cDNA in Clone ID NO:Z which encodes a mature polypeptide; (d) a nucleotide sequence in SEQ ID NO:X or the cDNA sequence of Clone ID NO:Z, which encodes a biologically active fragment of a

601607	Rhabdoid tumors
601649	Blepharophimosis, epicanthus inversus, and ptosis, type 2
601652	Glaucoma 1A, primary open angle, juvenile-onset, 137750
601680	Distal arthrogryposis, type 2B
601691	Retinitis pigmentosa-19, 601718
601691	Stargardt disease-1, 248200
601691	Cone-rod dystrophy 3
601691	Fundus flavimaculatus with macular dystrophy, 248200
601692	Reis-Bucklers corneal dystrophy
601692	Corneal dystrophy, Avellino type
601692	Corneal dystrophy, Groenouw type I, 121900
601692	Corneal dystrophy, lattice type I, 122200
601718	Retinitis pigmentosa-19
601780	Ceroid-lipofuscinosis, neuronal-6, variant late infantile
601800	[Hair color, brown]
601843	Hypothyroidism, congenital, 274400
601884	[High bone mass]
601969	Medulloblastoma, 155255
601969	Glioblastoma multiforme, 137800
602014	Hypomagnesemia with secondary hypocalcemia
602081	Speech-language disorder-1
602085	Postaxial polydactyly, type A2
602088	Nephronophthisis, infantile
602089	Hemangioma, capillary, hereditary
602091	Marfan syndrome, atypical
602092	Deafness, autosomal recessive 18
602094	Lipodystrophy, familial partial
602096	Alzheimer disease-5
602116	Glioma
602121	Deafness, autosomal dominant nonsyndromic sensorineural, 1, 124900
602134	Tremor, familial essential, 2
602136	Refsum disease, infantile, 266510
602136	Zellweger syndrome-1, 214100
602136	Adrenoleukodystrophy, neonatal, 202370
602225	Cone-rod retinal dystrophy-2, 120970
602225	Leber congenital amaurosis, type III
602447	Coronary artery disease, susceptibility to
602460	Deafness, autosomal dominant 15, 602459
602491	Hyperlipidemia, familial combined, 1
602574	Deafness, autosomal dominant 12, 601842
602574	Deafness, autosomal dominant 8, 601543
602629	Dystonia-6, torsion
602631	Rhabdomyosarcoma, 268210
602631	Breast Cancer

300067	Subcortical laminar heterotopia, X-linked dominant
300067	Lissencephaly, X-linked
300121	Subcortical laminar heterotopia, X-linked, 300067
300121	Lissencephaly, X-linked, 300067
300300	XLA and isolated growth hormone deficiency, 307200
300300	Agammaglobulinemia, type 1, X-linked
301201	Amelogenesis imperfecta-3, hypoplastic type
301835	Arts syndrome
309300	Megalocornea, X-linked
309605	Mental retardation, X-linked, syndromic-4, with congenital contractures and low fingertip arches
311850	Phosphoribosyl pyrophosphate synthetase-related gout
600035	Schizencephaly
600040	Colorectal cancer
600044	Thrombocythemia, essential, 187950
600045	Xeroderma pigmentosum, group E, subtype 2
600143	Epilepsy, progressive, with mental retardation
600224	Spinocerebellar ataxia-5
600228	Pseudohypoaldosteronism, type I, 264350
600234	HMG-CoA synthase-2 deficiency
600319	Diabetes mellitus, insulin-dependent, 4
600374	Bardet-Biedl syndrome 4
600414	Adrenoleukodystrophy, neonatal, 202370
600528	CPT deficiency, hepatic, type I, 255120
600542	Chondrosarcoma, extraskeletal myxoid
600618	Leukemia, acute lymphoblastic
600631	Enuresis, nocturnal, 1
600698	Salivary adenoma
600698	Uterine leiomyoma
600698	Lipoma
600698	Lipomatosis, multiple, 151900
600807	Bronchial asthma
600808	Enuresis, nocturnal, 2
600811	Xeroderma pigmentosum, group E, DDB-negative subtype, 278740
600850	Schizophrenia disorder-4
600856	Beckwith-Wiedemann syndrome, 130650
600882	Charcot-Marie-Tooth neuropathy-2B
600897	Cataract, zonular pulverulent-1, 116200
601105	Pycnodysostosis, 265800
601385	Prostate cancer
601412	Deafness, autosomal dominant 7
601414	Retinitis pigmentosa-18
601458	Inflammatory bowel disease-2
601472	Charcot-Marie-Tooth neuropathy-2D
601596	Charcot-Marie-Tooth neuropathy, demyelinating

230800	Gaucher disease
230800	Gaucher disease with cardiovascular calcification
231675	Glutaricaciduria, type IIC
231680	Glutaricaciduria, type IIA
232600	McArdle disease
234200	Neurodegeneration with brain iron accumulation
238600	Chylomicronemia syndrome, familial
238600	Combined hyperlipemia, familial
238600	Hyperlipoproteinemia I
238600	Lipoprotein lipase deficiency
243500	Isovalericacidemia
245200	Krabbe disease
246600	Pancreatic lipase deficiency
246900	Lipoamide dehydrogenase deficiency
250100	Metachromatic leukodystrophy
250800	Methemoglobinemia, type I
250800	Methemoglobinemia, type II
251600	Microphthalmia, autosomal recessive
253800	Walker-Warburg syndrome, 236670
253800	Fukuyama type congenital muscular dystrophy
254770	Epilepsy, juvenile myoclonic
256731	Ceroid-lipofuscinosis, neuronal-5, variant late infantile
258501	3-methylglutaconicaciduria, type III
258900	Oroticaciduria
259700	Osteopetrosis, recessive
259730	Renal tubular acidosis-osteopetrosis syndrome
259770	Osteoporosis-pseudoglioma syndrome
261640	Phenylketonuria due to PTS deficiency
263700	Porphyria, congenital erythropoietic
264300	Pseudohermaphroditism, male, with gynecomastia
266200	Anemia, hemolytic, due to PK deficiency
266300	[Hair color, red]
270100	Situs inversus viscerum
272800	Tay-Sachs disease
272800	[Hex A pseudodeficiency]
272800	GM2-gangliosidosis, juvenile, adult
274500	Thyroid iodine peroxidase deficiency
274500	Goiter, congenital
274500	Hypothyroidism, congenital
274600	Pendred syndrome
274600	Deafness, autosomal recessive 4
276700	Tyrosinemia, type I
276900	Usher syndrome, type 1A
277900	Wilson disease
278700	Xeroderma pigmentosum, group A
300046	Mental retardation, X-linked 23, nonspecific

180721	Retinitis pigmentosa, digenic
180840	Susceptibility to IDDM
181031	Oguchi disease-1, 258100
181430	Scapuloperoneal syndrome, myopathic type
181460	Schistosoma mansoni, susceptibility/resistance to
181600	Sclerolylosis
182600	Spastic paraplegia-3A
182860	Pyropoikilocytosis
182860	Spherocytosis, recessive
182860	Elliptocytosis-2
186855	Leukemia-2, T-cell acute lymphoblastic
186921	Leukemia, T-cell acute lymphoblastic
188400	Velocardiofacial syndrome, 192430
188400	DiGeorge syndrome
188540	Hypothyroidism, nongoitrous
189800	Preeclampsia/eclampsia
190020	Bladder cancer, 109800
190300	Tremor, familial essential, 1
190450	Hemolytic anemia due to triosephosphate isomerase deficiency
191181	Cervical carcinoma
191290	Segawa syndrome, recessive
191315	Insensitivity to pain, congenital, with anhidrosis, 256800
192340	Diabetes insipidus, neurohypophyseal, 125700
192500	Jervell and Lange-Nielsen syndrome, 220400
192500	Long QT syndrome-1
192974	Neonatal alloimmune thrombocytopenia
192974	Glycoprotein Ia deficiency
193235	Vitreoretinopathy, neovascular inflammatory
194071	Wilms tumor, type 2
194071	Adrenocortical carcinoma, hereditary, 202300
200990	Acrocallosal syndrome
203740	Alpha-ketoglutarate dehydrogenase deficiency
204500	Ceroid-lipofuscinosis, neuronal 2, classic late infantile
208400	Aspartylglucosaminuria
209901	Bardet-Biedl syndrome 1
210900	Bloom syndrome
216550	Cohen syndrome
216950	C1r/C1s deficiency, combined
217095	Conotruncal cardiac anomalies
218000	Andermann syndrome
222800	Hemolytic anemia due to bisphosphoglycerate mutase deficiency
223900	Dysautonomia, familial
227220	[Eye color, brown]
229600	Fructose intolerance

142680	Periodic fever, familial
145001	Hyperparathyroidism-jaw tumor syndrome
146200	Hypoparathyroidism, familial
146790	Lupus nephritis, susceptibility to
147050	Atopy
148370	Keratolytic winter erythema
151670	Hepatic lipase deficiency
152445	Vohwinkel syndrome, 124500
152445	Erythrokeratoderma, progressive symmetric, 602036
153455	Cutis laxa, recessive, type I, 219100
153700	Macular dystrophy, vitelliform type
154276	Malignant hyperthermia susceptibility 3
154400	Acrofacial dysostosis, Nager type
154550	Carbohydrate-deficient glycoprotein syndrome, type Ib, 602579
159000	Muscular dystrophy, limb-girdle, type 1A
159001	Muscular dystrophy, limb-girdle, type 1B
160900	Myotonic dystrophy
161015	Mitochondrial complex I deficiency, 252010
164009	Leukemia, acute promyelocytic, NUPA/RARA type
164860	Renal cell carcinoma, papillary, familial and sporadic
166800	Otosclerosis
168461	Multiple myeloma, 254250
168461	Parathyroid adenomatosis 1
168461	Centrocytic lymphoma
171650	Lysosomal acid phosphatase deficiency
173360	Thrombophilia due to excessive plasminogen activator inhibitor
173360	Hemorrhagic diathesis due to PAI1 deficiency
173850	Polio, susceptibility to
174000	Medullary cystic kidney disease, AD
176260	Episodic ataxia/myokymia syndrome, 160120
176730	Diabetes mellitus, rare form
176730	Hyperproinsulinemia, familial
176730	MODY, one form
176930	Dysprothrombinemia
176930	Hypoprothrombinemia
177400	Apnea, postanesthetic
179095	Male infertility
179755	Renal cell carcinoma, papillary, 1
180104	Retinitis pigmentosa-9
180105	Retinitis pigmentosa-10
180200	Osteosarcoma, 259500
180200	Pinealoma with bilateral retinoblastoma
180200	Retinoblastoma
180200	Bladder cancer, 109800

126452	Autonomic nervous system dysfunction
126452	[Novelty seeking personality]
126650	Chloride diarrhea, congenital, Finnish type, 214700
126650	Colon cancer
130410	Glutaricaciduria, type IIB
131100	Multiple endocrine neoplasia I
131100	Prolactinoma, hyperparathyroidism, carcinoid syndrome
131100	Carcinoid tumor of lung
131400	Eosinophilia, familial
131440	Eosinophilic myeloproliferative disorder
132800	Basal cell carcinoma
132800	Epithelioma, self-healing, squamous 1, Ferguson-Smith type
132810	Diphenylhydantoin toxicity
132810	Fetal hydantoin syndrome
133701	Exostoses, multiple, type 2
133780	Vitreoretinopathy, exudative, familial
134790	Hyperferritinemia-cataract syndrome, 600886
134820	Dysfibrinogenemia, alpha type, causing bleeding diathesis
134820	Dysfibrinogenemia, alpha type, causing recurrent thrombosis
134820	Amyloidosis, hereditary renal, 105200
134830	Dysfibrinogenemia, beta type
134850	Dysfibrinogenemia, gamma type
134850	Hypofibrinogenemia, gamma type
135940	Ichthyosis vulgaris, 146700
138040	Cortisol resistance
138079	Hyperinsulinism, familial, 602485
138079	MODY, type 2, 125851
138160	Diabetes mellitus, noninsulin-dependent
138160	Fanconi-Bickel syndrome, 227810
138570	Non-insulin dependent diabetes mellitus, susceptibility to
139130	Hypertension, essential, susceptibility to, 145500
139191	Growth hormone deficient dwarfism
141900	Methemoglobinemias, beta-
141900	Sickle cell anemia
141900	Thalassemias, beta-
141900	Erythremias, beta-
141900	HPFH, deletion type
141900	Heinz body anemias, beta-
142000	Thalassemia due to Hb Lepore
142000	Thalassemia, delta-
142200	HPFH, nondeletion type A
142250	HPFH, nondeletion type G
142270	Hereditary persistence of fetal hemoglobin
142360	Thrombophilia due to heparin cofactor II deficiency

						(Pharmacia) with a modified polylinker
L0809	NCI_CGAP_Pr28		prostate			pT7T3D-Pac (Pharmacia) with a modified polylinker
L2251	Human fetal lung	Fetal lung				
L5623	NCI_CGAP_Skn4	squamous cell carcinoma	skin			pCMV- SPORT6

TABLE 5

OMIM Reference	Description
102200	Somatotrophinoma
102540	Cardiomyopathy, idiopathic dilated
102578	Leukemia, acute promyelocytic, PML/RARA type
102770	Myoadenylate deaminase deficiency
103950	Emphysema due to alpha-2-macroglobulin deficiency
104170	NAGA deficiency, mild
104170	Schindler disease
104170	Kanzaki disease
104770	Amyloidosis, secondary, susceptibility to
106100	Angioedema, hereditary
107250	Anterior segment mesenchymal dysgenesis
107670	Apolipoprotein A-II deficiency
107776	Colton blood group, 110450
107970	Arrhythmogenic right ventricular dysplasia-1
108985	Atrophia areata
109400	Basal cell nevus syndrome
109700	Hemodialysis-related amyloidosis
110700	Vivax malaria, susceptibility to
113900	Heart block, progressive familial, type I
115200	Cardiomyopathy, dilated, 1A
115470	Cat eye syndrome
115650	Cataract, anterior polar-1
118485	Polycystic ovary syndrome with hyperandrogenemia
118511	Schizophrenia, neurophysiologic defect in
120580	C1r/C1s deficiency, combined
121050	Contractural arachnodactyly, congenital
123270	[Creatine kinase, brain type, ectopic expression of]
125852	Insulin-dependent diabetes mellitus-2
126340	Xeroderma pigmentosum, group D, 278730
126391	DNA ligase I deficiency

						modified polylinker
L0788	NCI_CGAP_Sub2					pT7T3D-Pac (Pharmacia) with a modified polylinker
L0789	NCI_CGAP_Sub3					pT7T3D-Pac (Pharmacia) with a modified polylinker
L0790	NCI_CGAP_Sub4					pT7T3D-Pac (Pharmacia) with a modified polylinker
L0791	NCI_CGAP_Sub5					pT7T3D-Pac (Pharmacia) with a modified polylinker
L0792	NCI_CGAP_Sub6					pT7T3D-Pac (Pharmacia) with a modified polylinker
L0794	NCI_CGAP_GC6	pooled germ cell tumors				pT7T3D-Pac (Pharmacia) with a modified polylinker
L0796	NCI_CGAP_Brn50	medulloblastoma	brain			pT7T3D-Pac (Pharmacia) with a modified polylinker
L0800	NCI_CGAP_Co16	colon tumor, RER+	colon			pT7T3D-Pac (Pharmacia) with a modified polylinker
L0803	NCI_CGAP_Kid11		kidney			pT7T3D-Pac (Pharmacia) with a modified polylinker
L0804	NCI_CGAP_Kid12	2 pooled tumors (clear cell type)	kidney			pT7T3D-Pac (Pharmacia) with a modified polylinker
L0805	NCI_CGAP_Lu24	carcinoid	lung			pT7T3D-Pac (Pharmacia) with a modified polylinker
L0806	NCI_CGAP_Lu19	squamous cell carcinoma, poorly differentiated (4	lung			pT7T3D-Pac (Pharmacia) with a modified polylinker
L0807	NCI_CGAP_Ov18	fibrotheoma	ovary			pT7T3D-Pac

L0773	NCI_CGAP_Co9	colon tumor RER+	colon			pT7T3D-Pac (Pharmacia) with a modified polylinker
L0774	NCI_CGAP_Kid3		kidney			pT7T3D-Pac (Pharmacia) with a modified polylinker
L0775	NCI_CGAP_Kid5	2 pooled tumors (clear cell type)	kidney			pT7T3D-Pac (Pharmacia) with a modified polylinker
L0776	NCI_CGAP_Lu5	carcinoid	lung			pT7T3D-Pac (Pharmacia) with a modified polylinker
L0777	Soares_NhHMPu_S1	Pooled human melanocyte, fetal heart, and pregnant	mixed (see below)			pT7T3D-Pac (Pharmacia) with a modified polylinker
L0779	Soares_NFL_T_GBC_S1		pooled			pT7T3D-Pac (Pharmacia) with a modified polylinker
L0780	Soares_NSF_F8_9W_OT_PA_P_S1		pooled			pT7T3D-Pac (Pharmacia) with a modified polylinker
L0782	NCI_CGAP_Pr21	normal prostate	prostate			pT7T3D-Pac (Pharmacia) with a modified polylinker
L0783	NCI_CGAP_Pr22	normal prostate	prostate			pT7T3D-Pac (Pharmacia) with a modified polylinker
L0784	NCI_CGAP_Lei2	leiomyosarcoma	soft tissue			pT7T3D-Pac (Pharmacia) with a modified polylinker
L0785	Barstead spleen HPLRB2		spleen			pT7T3D-Pac (Pharmacia) with a modified polylinker
L0786	Soares_NbHFB		whole brain			pT7T3D-Pac (Pharmacia) with a modified polylinker
L0787	NCI_CGAP_Sub1					pT7T3D-Pac (Pharmacia) with a

						with a modified polylinker
L0761	NCI_CGAP_CLL1	B-cell, chronic lymphocytic leukemia				pT7T3D-Pac (Pharmacia) with a modified polylinker
L0762	NCI_CGAP_Br1.1	breast				pT7T3D-Pac (Pharmacia) with a modified polylinker
L0763	NCI_CGAP_Br2	breast				pT7T3D-Pac (Pharmacia) with a modified polylinker
L0764	NCI_CGAP_Co3	colon				pT7T3D-Pac (Pharmacia) with a modified polylinker
L0765	NCI_CGAP_Co4	colon				pT7T3D-Pac (Pharmacia) with a modified polylinker
L0766	NCI_CGAP_GCB1	germinal center B cell				pT7T3D-Pac (Pharmacia) with a modified polylinker
L0767	NCI_CGAP_GC3	pooled germ cell tumors				pT7T3D-Pac (Pharmacia) with a modified polylinker
L0768	NCI_CGAP_GC4	pooled germ cell tumors				pT7T3D-Pac (Pharmacia) with a modified polylinker
L0769	NCI_CGAP_Brn25	anaplastic oligodendroglioma	brain			pT7T3D-Pac (Pharmacia) with a modified polylinker
L0770	NCI_CGAP_Brn23	glioblastoma (pooled)	brain			pT7T3D-Pac (Pharmacia) with a modified polylinker
L0771	NCI_CGAP_Co8	adenocarcinoma	colon			pT7T3D-Pac (Pharmacia) with a modified polylinker
L0772	NCI_CGAP_Co10	colon tumor RER+	colon			pT7T3D-Pac (Pharmacia) with a modified polylinker

	INFLS		Spleen			(Pharmacia) with a modified polylinker
L0749	Soares_fetal_liver_splee n_1NFLS_S1		Liver and Spleen			pT7T3D (Pharmacia) with a modified polylinker
L0750	Soares_fetal_lung_NbH L19W		lung			pT7T3D (Pharmacia) with a modified polylinker
L0751	Soares ovary tumor NbHOT	ovarian tumor	ovary			pT7T3D (Pharmacia) with a modified polylinker
L0752	Soares_parathyroid_tum or_NbHPA	parathyroid tumor	parathyroi d gland			pT7T3D (Pharmacia) with a modified polylinker
L0753	Soares_pineal_gland_N 3HPG		pineal gland			pT7T3D (Pharmacia) with a modified polylinker
L0754	Soares placenta Nb2HP		placenta			pT7T3D (Pharmacia) with a modified polylinker
L0755	Soares_placenta_8to9we eks_2NbHP8to9W		placenta			pT7T3D (Pharmacia) with a modified polylinker
L0756	Soares_multiple_scleros is_2NbHMSP	multiple sclerosis lesions				pT7T3D (Pharmacia) with a modified polylinker V_TYPE
L0757	Soares_senescent_fibro blasts_NbHSF	senescent fibroblast				pT7T3D (Pharmacia) with a modified polylinker V_TYPE
L0758	Soares_testis_NHT					pT7T3D-Pac (Pharmacia) with a modified polylinker
L0759	Soares_total_fetus_Nb2 HF8_9w					pT7T3D-Pac (Pharmacia) with a modified polylinker
L0760	Barstead aorta HPLRB3	aorta				pT7T3D-Pac (Pharmacia)

		with signet r				
L0663	NCI_CGAP_Ut2	moderately-differentiated endometrial adenocarcinoma	uterus			pCMV-SPORT6
L0664	NCI_CGAP_Ut3	poorly-differentiated endometrial adenocarcinoma,	uterus			pCMV-SPORT6
L0665	NCI_CGAP_Ut4	serous papillary carcinoma, high grade, 2 pooled t	uterus			pCMV-SPORT6
L0666	NCI_CGAP_Ut1	well-differentiated endometrial adenocarcinoma, 7	uterus			pCMV-SPORT6
L0667	NCI_CGAP_CML1	myeloid cells, 18 pooled CML cases, BCR/ABL rearra	whole blood			pCMV-SPORT6
L0717	Gessler Wilms tumor					pSPORT1
L0718	Testis 5					pSPORT1
L0731	Soares_pregnant_uterus_NbHPU		uterus			pT7T3-Pac
L0738	Human colorectal cancer					pT7T3D
L0740	Soares melanocyte 2NbHM	melanocyte				pT7T3D (Pharmacia) with a modified polylinker
L0741	Soares adult brain N2b4HB55Y		brain			pT7T3D (Pharmacia) with a modified polylinker
L0742	Soares adult brain N2b5HB55Y		brain			pT7T3D (Pharmacia) with a modified polylinker
L0743	Soares breast 2NbHBst		breast			pT7T3D (Pharmacia) with a modified polylinker
L0744	Soares breast 3NbHBst		breast			pT7T3D (Pharmacia) with a modified polylinker
L0745	Soares retina N2b4HR	retina	eye			pT7T3D (Pharmacia) with a modified polylinker
L0746	Soares retina N2b5HR	retina	eye			pT7T3D (Pharmacia) with a modified polylinker
L0747	Soares_fetal_heart_NbH H19W		heart			pT7T3D (Pharmacia) with a modified polylinker
L0748	Soares fetal liver spleen		Liver and			pT7T3D

						SPORT4
L0633	NCI_CGAP_Lu6	small cell carcinoma	lung			pCMV-SPORT4
L0636	NCI_CGAP_Pit1	four pooled pituitary adenomas	brain			pCMV-SPORT6
L0637	NCI_CGAP_Brn53	three pooled meningiomas	brain			pCMV-SPORT6
L0638	NCI_CGAP_Brn35	tumor, 5 pooled (see description)	brain			pCMV-SPORT6
L0640	NCI_CGAP_Br18	four pooled high-grade tumors, including two prima	breast			pCMV-SPORT6
L0641	NCI_CGAP_Co17	juvenile granulosa tumor	colon			pCMV-SPORT6
L0642	NCI_CGAP_Co18	moderately differentiated adenocarcinoma	colon			pCMV-SPORT6
L0643	NCI_CGAP_Co19	moderately differentiated adenocarcinoma	colon			pCMV-SPORT6
L0644	NCI_CGAP_Co20	moderately differentiated adenocarcinoma	colon			pCMV-SPORT6
L0646	NCI_CGAP_Co14	moderately-differentiated adenocarcinoma	colon			pCMV-SPORT6
L0647	NCI_CGAP_Sar4	five pooled sarcomas, including myxoid liposarcoma	connective tissue			pCMV-SPORT6
L0648	NCI_CGAP_Eso2	squamous cell carcinoma	esophagus			pCMV-SPORT6
L0649	NCI_CGAP_GU1	2 pooled high-grade transitional cell tumors	genitourinary tract			pCMV-SPORT6
L0650	NCI_CGAP_Kid13	2 pooled Wilms' tumors, one primary and one metast	kidney			pCMV-SPORT6
L0651	NCI_CGAP_Kid8	renal cell tumor	kidney			pCMV-SPORT6
L0652	NCI_CGAP_Lu27	four pooled poorly-differentiated adenocarcinomas	lung			pCMV-SPORT6
L0653	NCI_CGAP_Lu28	two pooled squamous cell carcinomas	lung			pCMV-SPORT6
L0655	NCI_CGAP_Lym12	lymphoma, follicular mixed small and large cell	lymph node			pCMV-SPORT6
L0656	NCI_CGAP_Ov38	normal epithelium	ovary			pCMV-SPORT6
L0657	NCI_CGAP_Ov23	tumor, 5 pooled (see description)	ovary			pCMV-SPORT6
L0658	NCI_CGAP_Ov35	tumor, 5 pooled (see description)	ovary			pCMV-SPORT6
L0659	NCI_CGAP_Pan1	adenocarcinoma	pancreas			pCMV-SPORT6
L0661	NCI_CGAP_Mel15	malignant melanoma, metastatic to lymph node	skin			pCMV-SPORT6
L0662	NCI_CGAP_Gas4	poorly differentiated adenocarcinoma	stomach			pCMV-SPORT6

L0581	Stratagene liver (#937224)		liver			pBluescript SK
L0586	HTCDL1					pBluescript SK(-)
L0588	Stratagene endothelial cell 937223					pBluescript SK-
L0589	Stratagene fetal retina 937202					pBluescript SK-
L0590	Stratagene fibroblast (#937212)					pBluescript SK-
L0591	Stratagene HeLa cell s3 937216					pBluescript SK-
L0592	Stratagene hNT neuron (#937233)					pBluescript SK-
L0593	Stratagene neuroepithelium (#937231)					pBluescript SK-
L0594	Stratagene neuroepithelium NT2RAMI 937234					pBluescript SK-
L0595	Stratagene NT2 neuronal precursor 937230	neuroepithelial cells	brain			pBluescript SK-
L0596	Stratagene colon (#937204)		colon			pBluescript SK-
L0597	Stratagene corneal stroma (#937222)		cornea			pBluescript SK-
L0598	Morton Fetal Cochlea	cochlea	ear			pBluescript SK-
L0599	Stratagene lung (#937210)		lung			pBluescript SK-
L0600	Weizmann Olfactory Epithelium	olfactory epithelium	nose			pBluescript SK-
L0601	Stratagene pancreas (#937208)		pancreas			pBluescript SK-
L0602	Pancreatic Islet	pancreatic islet	pancreas			pBluescript SK-
L0603	Stratagene placenta (#937225)		placenta			pBluescript SK-
L0604	Stratagene muscle 937209	muscle	skeletal muscle			pBluescript SK-
L0605	Stratagene fetal spleen (#937205)	fetal spleen	spleen			pBluescript SK-
L0606	NCI_CGAP_Lym5	follicular lymphoma	lymph node			pBluescript SK-
L0608	Stratagene lung carcinoma 937218	lung carcinoma	lung	NCI-H69		pBluescript SK-
L0612	Schiller oligodendroglioma	oligodendroglioma	brain			pBluescript SK- (Stratagene)
L0615	22 week old human fetal liver cDNA library					pBluescriptII SK(-)
L0622	HM1					pcDNAII (Invitrogen)
L0623	HM3	pectoral muscle (after mastectomy)				pcDNAII (Invitrogen)
L0629	NCI_CGAP_Mel3	metastatic melanoma to bowel	bowel (skin primary)			pCMV-SPORT4
L0631	NCI_CGAP_Br7		breast			pCMV-SPORT4
L0632	NCI_CGAP_Li5	hepatic adenoma	liver			pCMV-

L0499	NCI_CGAP_HSC2	stem cell 34+/38+	bone marrow			pAMP1
L0500	NCI_CGAP_Brn20	oligodendroglioma	brain			pAMP1
L0501	NCI_CGAP_Brn21	oligodendroglioma	brain			pAMP1
L0509	NCI_CGAP_Lu26	invasive adenocarcinoma	lung			pAMP1
L0512	NCI_CGAP_Ov36	borderline ovarian carcinoma	ovary			pAMP1
L0513	NCI_CGAP_Ov37	early stage papillary serous carcinoma	ovary			pAMP1
L0515	NCI_CGAP_Ov32	papillary serous carcinoma	ovary			pAMP1
L0516	Chromosome 19p12-p13.1 exon					pAMP10
L0517	NCI_CGAP_Pr1					pAMP10
L0518	NCI_CGAP_Pr2					pAMP10
L0519	NCI_CGAP_Pr3					pAMP10
L0520	NCI_CGAP_Alv1	alveolar rhabdomyosarcoma				pAMP10
L0521	NCI_CGAP_Ew1	Ewing's sarcoma				pAMP10
L0522	NCI_CGAP_Kid1	kidney				pAMP10
L0523	NCI_CGAP_Lip2	liposarcoma				pAMP10
L0524	NCI_CGAP_Li1	liver				pAMP10
L0526	NCI_CGAP_Pr12	metastatic prostate bone lesion				pAMP10
L0527	NCI_CGAP_Ov2	ovary				pAMP10
L0529	NCI_CGAP_Pr6	prostate				pAMP10
L0532	NCI_CGAP_Thy1	thyroid				pAMP10
L0533	NCI_CGAP_HSC1	stem cells	bone marrow			pAMP10
L0534	Chromosome 7 Fetal Brain cDNA Library	brain	brain			pAMP10
L0541	NCI_CGAP_Pr7	low-grade prostatic neoplasia	prostate			pAMP10
L0542	NCI_CGAP_Pr11	normal prostatic epithelial cells	prostate			pAMP10
L0543	NCI_CGAP_Pr9	normal prostatic epithelial cells	prostate			pAMP10
L0544	NCI_CGAP_Pr4	prostatic intraepithelial neoplasia - high grade	prostate			pAMP10
L0545	NCI_CGAP_Pr4.1	prostatic intraepithelial neoplasia - high grade	prostate			pAMP10
L0549	NCI_CGAP_HN10	carcinoma in situ from retromolar trigone				pAMP10
L0551	NCI_CGAP_HN7	normal squamous epithelium, floor of mouth				pAMP10
L0555	NCI_CGAP_Lu34	large cell carcinoma	lung			pAMP10
L0558	NCI_CGAP_Ov40	endometrioid ovarian metastasis	ovary			pAMP10
L0559	NCI_CGAP_Ov39	papillary serous ovarian metastasis	ovary			pAMP10
L0561	NCI_CGAP_HN11	normal squamous epithelium	tongue			pAMP10
L0564	Jia bone marrow stroma	bone marrow stroma				pBluescript
L0565	Normal Human Trabecular Bone Cells	Bone	Hip			pBluescript

L0375	NCI_CGAP_Kid6	kidney tumor	kidney			Bluescript SK-
L0376	NCI_CGAP_Lar1	larynx	larynx			Bluescript SK-
L0377	NCI_CGAP_HN2	squamous cell carcinoma from vocal cord	larynx			Bluescript SK-
L0378	NCI_CGAP_Lu1	lung tumor	lung			Bluescript SK-
L0379	NCI_CGAP_Lym3	lymphoma	lymph node			Bluescript SK-
L0381	NCI_CGAP_HN4	squamous cell carcinoma	pharynx			Bluescript SK-
L0382	NCI_CGAP_Pr25	epithelium (cell line)	prostate			Bluescript SK-
L0383	NCI_CGAP_Pr24	invasive tumor (cell line)	prostate			Bluescript SK-
L0384	NCI_CGAP_Pr23	prostate tumor	prostate			Bluescript SK-
L0385	NCI_CGAP_Gas1	gastric tumor	stomach			Bluescript SK-
L0387	NCI_CGAP_GCB0	germinal center B-cells	tonsil			Bluescript SK-
L0388	NCI_CGAP_HN6	normal gingiva (cell line from immortalized kerati				Bluescript SK-
L0389	NCI_CGAP_HN5	normal gingiva (cell line from primary keratinocyt				Bluescript SK-
L0415	b4HB3MA Cot8-HAP-Ft					Lafmid BA
L0435	Infant brain, LLNL array of Dr. M. Soares 1NIB					lafmid BA
L0438	normalized infant brain cDNA	total brain	brain			lafmid BA
L0439	Soares infant brain 1NIB		whole brain			Lafmid BA
L0443	b4HB3MK					Lafmid BK
L0451	N3HFLSK20					Lafmid K
L0455	Human retina cDNA randomly primed sublibrary	retina	eye			lambda gt10
L0456	Human retina cDNA Tsp509I-cleaved sublibrary	retina	eye			lambda gt10
L0462	WATM1					lambda gt11
L0471	Human fetal heart, Lambda ZAP Express					Lambda ZAP Express
L0480	Stratagene cat#937212 (1992)					Lambda ZAP, pBluescript SK(-)
L0483	Human pancreatic islet					Lambda ZAPII
L0485	STRATAGENE Human skeletal muscle cDNA library, cat. #936215.	skeletal muscle	leg muscle			Lambda ZAPII
L0497	NCI_CGAP_HSC4	CD34+, CD38- from normal bone marrow donor	bone marrow			pAMP1

T0110	Human colon carcinoma (HCC) cell line, remake					pBluescript SK-
T0115	Human Colon Carcinoma (HCC) cell line					pBluescript SK-
L0002	Atrium cDNA library Human heart					
L0004	ClonTech HL 1065a					
L0005	Clontech human aorta polyA+ mRNA (#6572)					
L0021	Human adult (K.Okubo)					
L0034	Human chromosome 14					
L0040	Human colon mucosa					
L0041	Human epidermal keratinocyte					
L0053	Human pancreatic tumor					
L0055	Human promyelocyte					
L0065	Liver HepG2 cell line.					
L0105	Human aorta polyA+ (TFujiwara)	aorta				
L0109	Human brain cDNA	brain				
L0118	Human fetal brain S. Meier-Ewert	brain				
L0142	Human placenta cDNA (TFujiwara)	placenta				
L0143	Human placenta polyA+ (TFujiwara)	placenta				
L0157	Human fetal brain (TFujiwara)		brain			
L0163	Human heart cDNA (YNakamura)		heart			
L0171	Human lung adenocarcinoma A549	lung adenocarcinoma		A549		
L0194	Human pancreatic cancer cell line Patu 8988t	pancreatic cancer		Patu 8988t		
L0351	Infant brain, Bento Soares					BA, M13-derived
L0352	Normalized infant brain, Bento Soares					BA, M13-derived
L0361	Stratagene ovary (#937217)		ovary			Bluescript SK
L0362	Stratagene ovarian cancer (#937219)					Bluescript SK-
L0363	NCI_CGAP_GC2	germ cell tumor				Bluescript SK-
L0364	NCI_CGAP_GC5	germ cell tumor				Bluescript SK-
L0366	Stratagene schizo brain S11	schizophrenic brain S-11 frontal lobe				Bluescript SK-
L0368	NCI_CGAP_SS1	synovial sarcoma				Bluescript SK-
L0369	NCI_CGAP_AA1	adrenal adenoma	adrenal gland			Bluescript SK-
L0371	NCI_CGAP_Br3	breast tumor	breast			Bluescript SK-
L0372	NCI_CGAP_Co12	colon tumor	colon			Bluescript SK-
L0373	NCI_CGAP_Co11	tumor	colon			Bluescript SK-
L0374	NCI_CGAP_Co2	tumor	colon			Bluescript SK-

S0460	Thyroid Tumour	Thyroid Tumour				pSport1
S0462	Thyroid Thyroiditis	Thyroid Thyroiditis				pSport1
S0464	Larynx Normal	Larynx Normal				pSport1
S0466	Larynx Tumor	Larynx Tumor			disease	pSport1
S0472	Lung Mesothelium	PYBT				pSport1
S0474	Human blood platelets	Platelets	Blood platelets			Other
S0665	Human Amygdala; re-excision	Amygdala				Uni-ZAP XR
S3012	Smooth Muscle Serum Treated, Norm	Smooth muscle	Pulmonary artery	Cell Line		pBluescript
S3014	Smooth muscle, serum induced, re-exc	Smooth muscle	Pulmonary artery	Cell Line		pBluescript
S6014	H. hypothalamus, frac A	Hypothalamus	Brain			ZAP Express
S6016	H. Frontal Cortex, Epileptic	H. Brain, Frontal Cortex, Epileptic	Brain		disease	Uni-ZAP XR
S6022	H. Adipose Tissue	Human Adipose Tissue				Uni-ZAP XR
S6024	Alzheimers, spongy change	Alzheimer's/Spongy change	Brain		disease	Uni-ZAP XR
S6026	Frontal Lobe, Dementia	Frontal Lobe dementia/Alzheimer's	Brain			Uni-ZAP XR
S6028	Human Manic Depression Tissue	Human Manic depression tissue	Brain		disease	Uni-ZAP XR
T0002	Activated T-cells	Activated T-Cell, PBL fraction	Blood	Cell Line		pBluescript SK-
T0003	Human Fetal Lung	Human Fetal Lung				pBluescript SK-
T0006	Human Pineal Gland	Human Pineal Gland				pBluescript SK-
T0008	Colorectal Tumor	Colorectal Tumor			disease	pBluescript SK-
T0010	Human Infant Brain	Human Infant Brain				Other
T0023	Human Pancreatic Carcinoma	Human Pancreatic Carcinoma			disease	pBluescript SK-
T0039	HSA 172 Cells	Human HSA172 cell line				pBluescript SK-
T0040	HSC172 cells	SA172 Cells				pBluescript SK-
T0041	Jurkat T-cell G1 phase	Jurkat T-cell				pBluescript SK-
T0042	Jurkat T-Cell, S phase	Jurkat T-Cell Line				pBluescript SK-
T0048	Human Aortic Endothelium	Human Aortic Endothelium				pBluescript SK-
T0049	Aorta endothelial cells + TNF-a	Aorta endothelial cells				pBluescript SK-
T0060	Human White Adipose	Human White Fat				pBluescript SK-
T0067	Human Thyroid	Human Thyroid				pBluescript SK-
T0069	Human Uterus, normal	Human Uterus, normal				pBluescript SK-
T0071	Human Bone Marrow	Human Bone Marrow				pBluescript SK-
T0082	Human Adult Retina	Human Adult Retina				pBluescript SK-
T0109	Human (HCC) cell line liver (mouse) metastasis, remake					pBluescript SK-

S0364	Human Quadriceps	Quadriceps muscle				pSport1
S0366	Human Soleus	Soleus Muscle				pSport1
S0368	Human Pancreatic Langerhans	Islets of Langerhans				pSport1
S0372	Larynx carcinoma III	Larynx carcinoma			disease	pSport1
S0374	Normal colon	Normal colon				pSport1
S0376	Colon Tumor	Colon Tumor			disease	pSport1
S0378	Pancreas normal PCA4 No	Pancreas Normal PCA4 No				pSport1
S0380	Pancreas Tumor PCA4 Tu	Pancreas Tumor PCA4 Tu			disease	pSport1
S0382	Larynx carcinoma IV	Larynx carcinoma			disease	pSport1
S0384	Tongue carcinoma	Tongue carcinoma			disease	pSport1
S0386	Human Whole Brain, re-excision	Whole brain	Brain			ZAP Express
S0388	Human Hypothalamus, schizophrenia, re-excision	Human Hypothalamus, Schizophrenia			disease	Uni-ZAP XR
S0390	Smooth muscle, control; re-excision	Smooth muscle	Pulmonary artery	Cell Line		Uni-ZAP XR
S0392	Salivary Gland	Salivary gland; normal				pSport1
S0394	Stomach; normal	Stomach; normal				pSport1
S0398	Testis; normal	Testis; normal				pSport1
S0402	Adrenal Gland, normal	Adrenal gland; normal				pSport1
S0404	Rectum normal	Rectum, normal				pSport1
S0406	Rectum tumour	Rectum tumour				pSport1
S0410	Colon, tumour	Colon, tumour				pSport1
S0412	Temporal cortex- Alzheimer; subtracted	Temporal cortex, alzheimer			disease	Other
S0418	CHME Cell Line; treated 5 hrs	CHME Cell Line; treated				pCMVSPORT 3.0
S0420	CHME Cell Line, untreated	CHME Cell line, untreated				pSport1
S0422	Mo7e Cell Line GM-CSF treated (1ng/ml)	Mo7e Cell Line GM-CSF treated (1ng/ml)				pCMVSPORT 3.0
S0424	TF-1 Cell Line GM-CSF Treated	TF-1 Cell Line GM-CSF Treated				pSport1
S0426	Monocyte activated; re-excision	Monocyte-activated	blood	Cell Line		Uni-ZAP XR
S0428	Neutrophils control; re-excision	human neutrophils	blood	Cell Line		Uni-ZAP XR
S0430	Aryepiglottis Normal	Aryepiglottis Normal				pSport1
S0432	Sinus piniformis Tumour	Sinus piniformis Tumour				pSport1
S0438	Liver Normal Met5No	Liver Normal Met5No				pSport1
S0440	Liver Tumour Met 5 Tu	Liver Tumour				pSport1
S0442	Colon Normal	Colon Normal				pSport1
S0444	Colon Tumor	Colon Tumour			disease	pSport1
S0446	Tongue Tumour	Tongue Tumour				pSport1
S0448	Larynx Normal	Larynx Normal				pSport1
S0450	Larynx Tumour	Larynx Tumour				pSport1
S0452	Thymus	Thymus				pSport1
S0454	Placenta	Placenta	Placenta			pSport1
S0456	Tongue Normal	Tongue Normal				pSport1
S0458	Thyroid Normal (SDCA2 No)	Thyroid normal				pSport1

		Pituitary)				
S0268	PRMIX	PRMIX (Human Prostate)	prostate			PCRII
S0270	PTMIX	PTMIX (Human Thymus)	Thymus			PCRII
S0274	PCMIX	PCMIX (Human Cerebellum)	Brain			PCRII
S0276	Synovial hypoxia-RSF subtracted	Synovial fobroblasts (rheumatoid)	Synovial tissue			pSport1
S0278	H Macrophage (GM-CSF treated), re-excision	Macrophage (GM-CSF treated)				Uni-ZAP XR
S0280	Human Adipose Tissue, re-excision	Human Adipose Tissue				Uni-ZAP XR
S0282	Brain Frontal Cortex, re-excision	Brain frontal cortex	Brain			Lambda ZAP II
S0298	Bone marrow stroma,treated	Bone marrow stroma,treatedSB	Bone marrow			pSport1
S0300	Frontal lobe,dementia;re-excision	Frontal Lobe dementia/Alzheimer's	Brain			Uni-ZAP XR
S0306	Larynx normal #10 261-273	Larynx normal				pSport1
S0308	Spleen/normal	Spleen normal				pSport1
S0310	Normal trachea	Normal trachea				pSport1
S0312	Human osteoarthritic;fraction II	Human osteoarthritic cartilage			disease	pSport1
S0314	Human osteoarthritic;fraction I	Human osteoarthritic cartilage			disease	pSport1
S0316	Human Normal Cartilage,Fraction I	Human Normal Cartilage				pSport1
S0318	Human Normal Cartilage Fraction II	Human Normal Cartilage				pSport1
S0322	Siebben Polyposis	Siebben Polyposis				pSport1
S0328	Palate carcinoma	Palate carcinoma	Uvula		disease	pSport1
S0330	Palate normal	Palate normal	Uvula			pSport1
S0332	Pharynx carcinoma	Pharynx carcinoma	Hypopharynx			pSport1
S0336	Human Normal Cartilage Fraction IV	Human Normal Cartilage				pSport1
S0338	Human Osteoarthritic Cartilage Fraction III	Human osteoarthritic cartilage			disease	pSport1
S0340	Human Osteoarthritic Cartilage Fraction IV	Human osteoarthritic cartilage			disease	pSport1
S0342	Adipocytes;re-excision	Human Adipocytes from Osteoclastoma				Uni-ZAP XR
S0344	Macrophage-oxLDL; re-excision	macrophage-oxidized LDL treated	blood	Cell Line		Uni-ZAP XR
S0346	Human Amygdala;re-excision	Amygdala				Uni-ZAP XR
S0350	Pharynx Carcinoma	Pharynx carcinoma	Hypopharynx		disease	pSport1
S0352	Larynx Carcinoma	Larynx carcinoma			disease	pSport1
S0354	Colon Normal II	Colon Normal	Colon			pSport1
S0356	Colon Carcinoma	Colon Carcinoma	Colon		disease	pSport1
S0358	Colon Normal III	Colon Normal	Colon			pSport1
S0360	Colon Tumor II	Colon Tumor	Colon		disease	pSport1

			marrow			ZAP II
S0140	eosinophil-IL5 induced	eosinophil	lung	Cell Line		Uni-ZAP XR
S0142	Macrophage-oxLDL	macrophage-oxidized LDL treated	blood	Cell Line		Uni-ZAP XR
S0144	Macrophage (GM-CSF treated)	Macrophage (GM-CSF treated)				Uni-ZAP XR
S0146	prostate-edited	prostate BPH	Prostate			Uni-ZAP XR
S0148	Normal Prostate	Prostate	prostate			Uni-ZAP XR
S0150	LNCAP prostate cell line	LNCAP Cell Line	Prostate	Cell Line		Uni-ZAP XR
S0152	PC3 Prostate cell line	PC3 prostate cell line				Uni-ZAP XR
S0182	Human B Cell 8866	Human B- Cell 8866				Uni-ZAP XR
S0190	Prostate BPH,Lib 2, subtracted	Human Prostate BPH				pSport1
S0192	Synovial Fibroblasts (control)	Synovial Fibroblasts				pSport1
S0194	Synovial hypoxia	Synovial Fibroblasts				pSport1
S0196	Synovial IL-1/TNF stimulated	Synovial Fibroblasts				pSport1
S0198	7TM-pbfd	PBLS, 7TM receptor enriched				PCRII
S0202	7TM-pbdd	PBLS, 7TM receptor enriched				PCRII
S0206	Smooth Muscle-HASTE normalized	Smooth muscle	Pulmonary artery	Cell Line		pBluescript
S0208	Messangial cell, frac 1	Messangial cell				pSport1
S0210	Messangial cell, frac 2	Messangial cell				pSport1
S0212	Bone Marrow Stromal Cell, untreated	Bone Marrow Stromal Cell, untreated				pSport1
S0214	Human Osteoclastoma, re-excision	Osteoclastoma	bone		disease	Uni-ZAP XR
S0216	Neutrophils IL-1 and LPS induced	human neutrophil induced	blood	Cell Line		Uni-ZAP XR
S0218	Apoptotic T-cell, re-excision	apoptotic cells		Cell Line		Uni-ZAP XR
S0222	H. Frontal cortex, epileptic; re-excision	H. Brain, Frontal Cortex, Epileptic	Brain		disease	Uni-ZAP XR
S0228	PSMIX	PBLS, 7TM receptor enriched				PCRII
S0242	Synovial Fibroblasts (II/TNF), subt	Synovial Fibroblasts				pSport1
S0250	Human Osteoblasts II	Human Osteoblasts	Femur		disease	pCMVSPORT 2.0
S0252	7TM-PIMIX	PBLS, 7TM receptor enriched				PCRII
S0254	7TM-PAMIX	PBLS, 7TM receptor enriched				PCRII
S0256	7TM-PHMIX	PBLS, 7TM receptor enriched				PCRII
S0258	7TM-PNMIX	PBLS, 7TM receptor enriched				PCRII
S0260	Spinal Cord, re-excision	Spinal cord	spinal cord			Uni-ZAP XR
S0264	PPMIX	PPMIX (Human	Pituitary			PCRII

	unamplified					
S0026	Stromal cell TF274	stromal cell	Bone marrow	Cell Line		Uni-ZAP XR
S0027	Smooth muscle, serum treated	Smooth muscle	Pulmonary artery	Cell Line		Uni-ZAP XR
S0028	Smooth muscle, control	Smooth muscle	Pulmonary artery	Cell Line		Uni-ZAP XR
S0029	brain stem	Brain stem	brain			Uni-ZAP XR
S0030	Brain pons	Brain Pons	Brain			Uni-ZAP XR
S0031	Spinal cord	Spinal cord	spinal cord			Uni-ZAP XR
S0032	Smooth muscle-IL1b induced	Smooth muscle	Pulmonary artery	Cell Line		Uni-ZAP XR
S0036	Human Substantia Nigra	Human Substantia Nigra				Uni-ZAP XR
S0037	Smooth muscle, IL1b induced	Smooth muscle	Pulmonary artery	Cell Line		Uni-ZAP XR
S0038	Human Whole Brain #2 - Oligo dT > 1.5Kb	Human Whole Brain #2				ZAP Express
S0039	Hypothalamus	Hypothalamus	Brain			Uni-ZAP XR
S0040	Adipocytes	Human Adipocytes from Osteoclastoma				Uni-ZAP XR
S0044	Prostate BPH	prostate BPH	Prostate		disease	Uni-ZAP XR
S0045	Endothelial cells-control	Endothelial cell	endothelial cell-lung	Cell Line		Uni-ZAP XR
S0046	Endothelial-induced	Endothelial cell	endothelial cell-lung	Cell Line		Uni-ZAP XR
S0049	Human Brain, Striatum	Human Brain, Striatum				Uni-ZAP XR
S0050	Human Frontal Cortex, Schizophrenia	Human Frontal Cortex, Schizophrenia			disease	Uni-ZAP XR
S0051	Human Hypothalamus, Schizophrenia	Human Hypothalamus, Schizophrenia			disease	Uni-ZAP XR
S0052	neutrophils control	human neutrophils	blood	Cell Line		Uni-ZAP XR
S0053	Neutrophils IL-1 and LPS induced	human neutrophil induced	blood	Cell Line		Uni-ZAP XR
S0106	STRIATUM DEPRESSION		BRAIN		disease	Uni-ZAP XR
S0110	Brain Amygdala Depression		Brain		disease	Uni-ZAP XR
S0112	Hypothalamus		Brain			Uni-ZAP XR
S0114	Anergic T-cell	Anergic T-cell		Cell Line		Uni-ZAP XR
S0116	Bone marrow	Bone marrow	Bone marrow			Uni-ZAP XR
S0122	Osteoclastoma-normalized A	Osteoclastoma	bone		disease	pBluescript
S0126	Osteoblasts	Osteoblasts	Knee	Cell Line		Uni-ZAP XR
S0132	Epithelial-TNF α and INF induced	Airway Epithelial				Uni-ZAP XR
S0134	Apoptotic T-cell	apoptotic cells		Cell Line		Uni-ZAP XR
S0136	PERM TF274	stromal cell	Bone	Cell Line		Lambda

	(9808C064R)	9808C064R				3.0
H0676	Colon, Cancer: (9808C064R)-total RNA	Colon Cancer 9808C064R				pCMVSPORT 3.0
H0677	TNFR degenerate oligo	B-Cells				PCR II
H0679	screened clones from Tonsil library	Human Tonsils				Other
H0682	Serous Papillary Adenocarcinoma	serous papillary adenocarcinoma (9606G304SPA3B)				pCMVSPORT 3.0
H0683	Ovarian Serous Papillary Adenocarcinoma	Serous papillary adenocarcinoma, stage 3C (9804G01)				pCMVSPORT 3.0
H0684	Serous Papillary Adenocarcinoma	Ovarian Cancer- 9810G606	Ovaries			pCMVSPORT 3.0
H0685	Adenocarcinoma of Ovary, Human Cell Line, # OVCAR-3	Adenocarcinoma of Ovary, Human Cell Line, # OVCAR-				pCMVSPORT 3.0
H0686	Adenocarcinoma of Ovary, Human Cell Line	Adenocarcinoma of Ovary, Human Cell Line, # SW-626				pCMVSPORT 3.0
H0687	Human normal ovary(#9610G215)	Human normal ovary(#9610G215)	Ovary			pCMVSPORT 3.0
H0688	Human Ovarian Cancer(#9807G017)	Human Ovarian cancer(#9807G017), mRNA from Maura Ru				pCMVSPORT 3.0
H0689	Ovarian Cancer	Ovarian Cancer, #9806G019				pCMVSPORT 3.0
H0690	Ovarian Cancer, # 9702G001	Ovarian Cancer, #9702G001				pCMVSPORT 3.0
H0691	Normal Ovary, #9710G208	normal ovary, #9710G208				pCMVSPORT 3.0
H0693	Normal Prostate #ODQ3958EN	Normal Prostate Tissue # ODQ3958EN				pCMVSPORT 3.0
H0694	Prostate gland adenocarcinoma	Prostate gland, adenocarcinoma, mod/diff, gleason	prostate gland			pCMVSPORT 3.0
H0695	mononucleocytes from patient	mononucleocytes from patient at Shady Grove Hospit				pCMVSPORT 3.0
N0006	Human Fetal Brain	Human Fetal Brain				
S0001	Brain frontal cortex	Brain frontal cortex	Brain			Lambda ZAP II
S0002	Monocyte activated	Monocyte-activated	blood	Cell Line		Uni-ZAP XR
S0003	Human Osteoclastoma	Osteoclastoma	bone		disease	Uni-ZAP XR
S0007	Early Stage Human Brain	Human Fetal Brain				Uni-ZAP XR
S0010	Human Amygdala	Amygdala				Uni-ZAP XR
S0011	STROMAL - OSTEOCLASTOMA	Osteoclastoma	bone		disease	Uni-ZAP XR
S0013	Prostate	Prostate	prostate			Uni-ZAP XR
S0016	Kidney Pyramids	Kidney pyramids	Kidney			Uni-ZAP XR
S0021	Whole brain	Whole brain	Brain			ZAP Express
S0022	Human Osteoclastoma Stromal Cells -	Osteoclastoma Stromal Cells				Uni-ZAP XR

	Diff. Adenocarcinoma, Metastatic	adenocarcinoma				
H0648	Ovary, Cancer: (4004562 B6) Papillary Serous Cystic Neoplasm, Low Malignant Pot	Papillary Cstic neoplasm of low malignant potentia			disease	pSport1
H0649	Lung, Normal: (4005313 B1)	Normal Lung				pSport1
H0650	B-Cells	B-Cells				pCMVSPORT 3.0
H0651	Ovary, Normal: (9805C040R)	Normal Ovary				pSport1
H0652	Lung, Normal: (4005313 B1)	Normal Lung				pSport1
H0653	Stromal Cells	Stromal Cells				pSport1
H0656	B-cells (unstimulated)	B-cells (unstimulated)				pSport1
H0657	B-cells (stimulated)	B-cells (stimulated)				pSport1
H0658	Ovary, Cancer (9809C332): Poorly differentiated adenocarcinoma	9809C332- Poorly differentiate	Ovary & Fallopian Tubes		disease	pSport1
H0659	Ovary, Cancer (15395A1F): Grade II Papillary Carcinoma	Grade II Papillary Carcinoma, Ovary	Ovary		disease	pSport1
H0660	Ovary, Cancer: (15799A1F) Poorly differentiated carcinoma	Poorly differentiated carcinoma, ovary			disease	pSport1
H0661	Breast, Cancer: (4004943 A5)	Breast cancer			disease	pSport1
H0662	Breast, Normal: (4005522B2)	Normal Breast - #4005522(B2)	Breast			pSport1
H0663	Breast, Cancer: (4005522 A2)	Breast Cancer - #4005522(A2)	Breast		disease	pSport1
H0664	Breast, Cancer: (9806C012R)	Breast Cancer	Breast		disease	pSport1
H0665	Stromal cells 3.88	Stromal cells 3.88				pSport1
H0666	Ovary, Cancer: (4004332 A2)	Ovarian Cancer, Sample #4004332A2			disease	pSport1
H0667	Stromal cells(HBM3.18)	Stromal cell(HBM 3.18)				pSport1
H0668	stromal cell clone 2.5	stromal cell clone 2.5				pSport1
H0669	Breast, Cancer: (4005385 A2)	Breast Cancer (4005385A2)	Breast			pSport1
H0670	Ovary, Cancer(4004650 A3): Well-Differentiated Micropapillary Serous Carcinoma	Ovarian Cancer - 4004650A3				pSport1
H0671	Breast, Cancer: (9802C02OE)	Breast Cancer-Sample # 9802C02OE				pSport1
H0672	Ovary, Cancer: (4004576 A8)	Ovarian Cancer(4004576A8)	Ovary			pSport1
H0673	Human Prostate Cancer, Stage B2; re-excision	Human Prostate Cancer, stage B2	Prostate			Uni-ZAP XR
H0674	Human Prostate Cancer, Stage C; re-excision	Human Prostate Cancer, stage C	Prostate			Uni-ZAP XR
H0675	Colon, Cancer:	Colon Cancer				pCMVSPORT

H0606	Human Primary Breast Cancer; re-excision	Human Primary Breast Cancer	Breast		disease	Uni-ZAP XR
H0610	H. Leukocytes, normalized cot 5A	H. Leukocytes				pCMVSPORT 1
H0615	Human Ovarian Cancer Reexcision	Ovarian Cancer	Ovary		disease	Uni-ZAP XR
H0616	Human Testes, Reexcision	Human Testes	Testis			Uni-ZAP XR
H0617	Human Primary Breast Cancer Reexcision	Human Primary Breast Cancer	Breast		disease	Uni-ZAP XR
H0618	Human Adult Testes, Large Inserts, Reexcision	Human Adult Testis	Testis			Uni-ZAP XR
H0619	Fetal Heart	Human Fetal Heart	Heart			Uni-ZAP XR
H0620	Human Fetal Kidney; Reexcision	Human Fetal Kidney	Kidney			Uni-ZAP XR
H0622	Human Pancreas Tumor; Reexcision	Human Pancreas Tumor	Pancreas		disease	Uni-ZAP XR
H0623	Human Umbilical Vein; Reexcision	Human Umbilical Vein Endothelial Cells	Umbilical vein			Uni-ZAP XR
H0624	12 Week Early Stage Human II; Reexcision	Twelve Week Old Early Stage Human	Embryo			Uni-ZAP XR
H0625	Ku 812F Basophils Line	Ku 812F Basophils				pSport1
H0626	Sa0s2 Cells; Untreated	Sa0s2 Cell Line; Untreated				pSport1
H0628	Human Pre-Differentiated Adipocytes	Human Pre-Differentiated Adipocytes				Uni-ZAP XR
H0631	Sa0s2, Dexamethosone Treated	Sa0s2 Cell Line; Dexamethosone Treated				pSport1
H0632	Hepatocellular Tumor; re-excision	Hepatocellular Tumor	Liver			Lambda ZAP II
H0633	Lung Carcinoma A549 TNFalpha activated	TNFalpha activated A549-Lung Carcinoma			disease	pSport1
H0634	Human Testes Tumor, re-excision	Human Testes Tumor	Testis		disease	Uni-ZAP XR
H0635	Human Activated T-Cells, re-excision	Activated T-Cells	Blood	Cell Line		Uni-ZAP XR
H0637	Dendritic Cells From CD34 Cells	Dendritic cells from CD34 cells				pSport1
H0638	CD40 activated monocyte dendritic cells	CD40 activated monocyte dendritic cells				pSport1
H0641	LPS activated derived dendritic cells	LPS activated monocyte derived dendritic cells				pSport1
H0642	Hep G2 Cells, lambda library	Hep G2 Cells				Other
H0644	Human Placenta (re-excision)	Human Placenta	Placenta			Uni-ZAP XR
H0645	Fetal Heart, re-excision	Human Fetal Heart	Heart			Uni-ZAP XR
H0646	Lung, Cancer (4005313 A3): Invasive Poorly Differentiated Lung Adenocarcinoma,	Metastatic squamous cell lung carcinoma, poorly di				pSport1
H0647	Lung, Cancer (4005163 B7): Invasive, Poorly	Invasive poorly differentiated lung			disease	pSport1

						3.0
H0561	L428	L428				pCMVSPORT 3.0
H0562	Human Fetal Brain, normalized c5-11-26	Human Fetal Brain				pCMVSPORT 2.0
H0570	Human Fetal Brain, normalized C500H	Human Fetal Brain				pCMVSPORT 2.0
H0571	Human Fetal Brain, normalized C500HE	Human Fetal Brain				pCMVSPORT 2.0
H0572	Human Fetal Brain, normalized AC5002	Human Fetal Brain				pCMVSPORT 2.0
H0574	Hepatocellular Tumor; re-excision	Hepatocellular Tumor	Liver		disease	Lambda ZAP II
H0575	Human Adult Pulmonary;re-excision	Human Adult Pulmonary	Lung			Uni-ZAP XR
H0576	Resting T-Cell; re- excision	T-Cells	Blood	Cell Line		Lambda ZAP II
H0580	Dendritic cells, pooled	Pooled dendritic cells				pCMVSPORT 3.0
H0581	Human Bone Marrow, treated	Human Bone Marrow	Bone Marrow			pCMVSPORT 3.0
H0583	B Cell lymphoma	B Cell Lymphoma	B Cell		disease	pCMVSPORT 3.0
H0584	Activated T-cells, 24 hrs, re-excision	Activated T-Cells	Blood	Cell Line		Uni-ZAP XR
H0586	Healing groin wound, 6.5 hours post incision	healing groin wound, 6.5 hours post incision - 2/	groin		disease	pCMVSPORT 3.0
H0587	Healing groin wound; 7.5 hours post incision	Groin-2/19/97	groin		disease	pCMVSPORT 3.0
H0589	CD34 positive cells (cord blood), re-ex	CD34 Positive Cells	Cord Blood			ZAP Express
H0590	Human adult small intestine, re-excision	Human Adult Small Intestine	Small Int.			Uni-ZAP XR
H0591	Human T-cell lymphoma; re-excision	T-Cell Lymphoma	T-Cell		disease	Uni-ZAP XR
H0592	Healing groin wound - zero hr post-incision (control)	HGS wound healing project; abdomen			disease	pCMVSPORT 3.0
H0593	Olfactory epithelium; nasal cavity	Olfactory epithelium from roof of left nasal cavity				pCMVSPORT 3.0
H0594	Human Lung Cancer; re- excision	Human Lung Cancer	Lung		disease	Lambda ZAP II
H0595	Stomach cancer (human); re-excision	Stomach Cancer - 5383A (human)			disease	Uni-ZAP XR
H0596	Human Colon Cancer; re-excision	Human Colon Cancer	Colon			Lambda ZAP II
H0597	Human Colon; re- excision	Human Colon				Lambda ZAP II
H0598	Human Stomach; re- excision	Human Stomach	Stomach			Uni-ZAP XR
H0599	Human Adult Heart; re- excision	Human Adult Heart	Heart			Uni-ZAP XR
H0600	Healing Abdomen wound; 70&90 min post incision	Abdomen			disease	pCMVSPORT 3.0
H0601	Healing Abdomen Wound; 15 days post incision	Abdomen			disease	pCMVSPORT 3.0
H0604	Human Pituitary, re- excision	Human Pituitary				pBluescript

H0510	Human Liver, normal	Human Liver, normal, Patient # 8	Liver			pCMVSPORT 3.0
H0518	pBMC stimulated w/ poly I/C	pBMC stimulated with poly I/C				pCMVSPORT 3.0
H0519	NTERA2, control	NTERA2, Teratocarcinoma cell line				pCMVSPORT 3.0
H0520	NTERA2 + retinoic acid, 14 days	NTERA2, Teratocarcinoma cell line				pSPORT1
H0521	Primary Dendritic Cells, lib 1	Primary Dendritic cells				pCMVSPORT 3.0
H0522	Primary Dendritic cells, frac 2	Primary Dendritic cells				pCMVSPORT 3.0
H0525	PCR, pBMC I/C treated	pBMC stimulated with poly I/C				PCRII
H0528	Poly[I]/Poly[C] Normal Lung Fibroblasts	Poly[I]/Poly[C] Normal Lung Fibroblasts				pCMVSPORT 3.0
H0529	Myeloid Progenitor Cell Line	TF-1 Cell Line; Myeloid progenitor cell line				pCMVSPORT 3.0
H0530	Human Dermal Endothelial Cells, untreated	Human Dermal Endothelial Cells; untreated				pSPORT1
H0538	Merkel Cells	Merkel cells	Lymph node			pSPORT1
H0539	Pancreas Islet Cell Tumor	Pancreas Islet Cell Tumour	Pancreas		disease	pSPORT1
H0542	T Cell helper I	Helper T cell				pCMVSPORT 3.0
H0543	T cell helper II	Helper T cell				pCMVSPORT 3.0
H0544	Human endometrial stromal cells	Human endometrial stromal cells				pCMVSPORT 3.0
H0545	Human endometrial stromal cells-treated with progesterone	Human endometrial stromal cells-treated with proge				pCMVSPORT 3.0
H0546	Human endometrial stromal cells-treated with estradiol	Human endometrial stromal cells-treated with estra				pCMVSPORT 3.0
H0547	NTERA2 teratocarcinoma cell line+retinoic acid (14 days)	NTERA2, Teratocarcinoma cell line				pSPORT1
H0549	H. Epididymus, caput & corpus	Human Epididymus, caput and corpus				Uni-ZAP XR
H0550	H. Epididymus, cauda	Human Epididymus, cauda				Uni-ZAP XR
H0551	Human Thymus Stromal Cells	Human Thymus Stromal Cells				pCMVSPORT 3.0
H0553	Human Placenta	Human Placenta				pCMVSPORT 3.0
H0555	Rejected Kidney, lib 4	Human Rejected Kidney	Kidney		disease	pCMVSPORT 3.0
H0556	Activated T-cell(12h)/Thiouridine-re-excision	T-Cells	Blood	Cell Line		Uni-ZAP XR
H0559	HL-60, PMA 4H, re-excision	HL-60 Cells, PMA stimulated 4H	Blood	Cell Line		Uni-ZAP XR
H0560	KMH2	KMH2				pCMVSPORT

		hiplipoma				
H0428	Human Ovary	Human Ovary Tumor	Ovary			pSport1
H0429	K562 + PMA (36 hrs), re-excision	K562 Cell line	cell line	Cell Line		ZAP Express
H0431	H. Kidney Medulla, re-excision	Kidney medulla	Kidney			pBluescript
H0433	Human Umbilical Vein Endothelial cells, frac B, re-excision	HUVE Cells	Umbilical vein	Cell Line		pBluescript
H0434	Human Brain, striatum, re-excision	Human Brain, Striatum				pBluescript
H0435	Ovarian Tumor 10-3-95	Ovarian Tumor, OV350721	Ovary			pCMVSPORT 2.0
H0436	Resting T-Cell Library, II	T-Cells	Blood	Cell Line		pSport1
H0437	H Umbilical Vein Endothelial Cells, frac A, re-excision	HUVE Cells	Umbilical vein	Cell Line		Lambda ZAP II
H0438	H. Whole Brain #2, re-excision	Human Whole Brain #2				ZAP Express
H0439	Human Eosinophils	Eosinophils				pBluescript
H0441	H. Kidney Cortex, subtracted	Kidney cortex	Kidney			pBluescript
H0442	H. Striatum Depression, subt II	Human Brain, Striatum Depression	Brain			pBluescript
H0444	Spleen metastatic melanoma	Spleen, Metastatic malignant melanoma	Spleen		disease	pSport1
H0445	Spleen, Chronic lymphocytic leukemia	Human Spleen, CLL	Spleen		disease	pSport1
H0453	H. Kidney Pyramid, subtracted	Kidney pyramids	Kidney			pBluescript
H0455	H. Striatum Depression, subt	Human Brain, Striatum Depression	Brain			pBluescript
H0457	Human Eosinophils	Human Eosinophils				pSport1
H0477	Human Tonsil, Lib 3	Human Tonsil	Tonsil			pSport1
H0478	Salivary Gland, Lib 2	Human Salivary Gland	Salivary gland			pSport1
H0483	Breast Cancer cell line, MDA 36	Breast Cancer Cell line, MDA 36				pSport1
H0484	Breast Cancer Cell line, angiogenic	Breast Cancer Cell line, Angiogenic, 36T3				pSport1
H0485	Hodgkin's Lymphoma I	Hodgkin's Lymphoma I			disease	pCMVSPORT 2.0
H0486	Hodgkin's Lymphoma II	Hodgkin's Lymphoma II			disease	pCMVSPORT 2.0
H0487	Human Tonsils, lib I	Human Tonsils				pCMVSPORT 2.0
H0488	Human Tonsils, Lib 2	Human Tonsils				pCMVSPORT 2.0
H0489	Crohn's Disease	Ileum	Intestine		disease	pSport1
H0492	HL-60, RA 4h, Subtracted	HL-60 Cells, RA stimulated for 4H	Blood	Cell Line		Uni-ZAP XR
H0494	Keratinocyte	Keratinocyte				pCMVSPORT 2.0
H0497	HEL cell line	HEL cell line		HEL 92.1.7		pSport1
H0506	Ulcerative Colitis	Colon	Colon			pSport1
H0509	Liver, Hepatoma	Human Liver, Hepatoma, patient 8	Liver		disease	pCMVSPORT 3.0

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H0354	Human Leukocytes	Human Leukocytes	Blood	Cell Line		pCMVSPORT 1
H0355	Human Liver	Human Liver, normal Adult				pCMVSPORT 1
H0356	Human Kidney	Human Kidney	Kidney			pCMVSPORT 1
H0357	H. Normalized Fetal Liver, II	Human Fetal Liver	Liver			Uni-ZAP XR
H0363	Human Brain Medulla, subtracted	Human Brain Medulla				pBluescript
H0369	H. Atrophic Endometrium	Atrophic Endometrium and myometrium				Uni-ZAP XR
H0370	H. Lymph node breast Cancer	Lymph node with Met. Breast Cancer			disease	Uni-ZAP XR
H0373	Human Heart	Human Adult Heart	Heart			pCMVSPORT 1
H0375	Human Lung	Human Lung				pCMVSPORT 1
H0379	Human Tongue, frac 1	Human Tongue				pSport1
H0380	Human Tongue, frac 2	Human Tongue				pSport1
H0381	Bone Cancer	Bone Cancer			disease	Uni-ZAP XR
H0386	Leukocyte and Lung; 4 screens	Human Leukocytes	Blood	Cell Line		pCMVSPORT 1
H0390	Human Amygdala Depression, re-excision	Human Amygdala Depression			disease	pBluescript
H0391	H. Meningioma, M6	Human Meningioma	brain			pSport1
H0392	H. Meningioma, M1	Human Meningioma	brain			pSport1
H0393	Fetal Liver, subtraction II	Human Fetal Liver	Liver			pBluescript
H0394	A-14 cell line	Redd-Sternberg cell				ZAP Express
H0396	L1 Cell line	Redd-Sternberg cell				ZAP Express
H0402	CD34 depleted Buffy Coat (Cord Blood), re-excision	CD34 Depleted Buffy Coat (Cord Blood)	Cord Blood			ZAP Express
H0408	Human kidney Cortex, subtracted	Human Kidney Cortex				pBluescript
H0411	H Female Bladder, Adult	Human Female Adult Bladder	Bladder			pSport1
H0412	Human umbilical vein endothelial cells, IL-4 induced	HUVE Cells	Umbilical vein	Cell Line		pSport1
H0413	Human Umbilical Vein Endothelial Cells, uninduced	HUVE Cells	Umbilical vein	Cell Line		pSport1
H0415	H. Ovarian Tumor, II, OV5232	Ovarian Tumor, OV5232	Ovary		disease	pCMVSPORT 2.0
H0416	Human Neutrophils, Activated, re-excision	Human Neutrophil - Activated	Blood	Cell Line		pBluescript
H0418	Human Pituitary, subtracted VII	Human Pituitary				pBluescript
H0421	Human Bone Marrow, re-excision	Bone Marrow				pBluescript
H0422	T-Cell PHA 16 hrs	T-Cells	Blood	Cell Line		pSport1
H0423	T-Cell PHA 24 hrs	T-Cells	Blood	Cell Line		pSport1
H0424	Human Pituitary, subt IX	Human Pituitary				pBluescript
H0427	Human Adipose	Human Adipose, left				pSport1

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H0269	Human Umbilical Vein Endothelial Cells, fract. B	HUVE Cells	Umbilical vein	Cell Line		Lambda ZAP II
H0270	HPAS (human pancreas, subtracted)	Human Pancreas	Pancreas			Uni-ZAP XR
H0271	Human Neutrophil, Activated	Human Neutrophil - Activated	Blood	Cell Line		Uni-ZAP XR
H0272	HUMAN TONSILS, FRACTION 2	Human Tonsil	Tonsil			Uni-ZAP XR
H0279	K562 cells	K562 Cell line	cell line	Cell Line		ZAP Express
H0280	K562 + PMA (36 hrs)	K562 Cell line	cell line	Cell Line		ZAP Express
H0286	Human OB MG63 treated (10 nM E2) fraction I	Human Osteoblastoma MG63 cell line	Bone	Cell Line		Uni-ZAP XR
H0288	Human OB HOS control fraction I	Human Osteoblastoma HOS cell line	Bone	Cell Line		Uni-ZAP XR
H0292	Human OB HOS treated (10 nM E2) fraction I	Human Osteoblastoma HOS cell line	Bone	Cell Line		Uni-ZAP XR
H0294	Amniotic Cells - TNF induced	Amniotic Cells - TNF induced	Placenta	Cell Line		Uni-ZAP XR
H0295	Amniotic Cells - Primary Culture	Amniotic Cells - Primary Culture	Placenta	Cell Line		Uni-ZAP XR
H0305	CD34 positive cells (Cord Blood)	CD34 Positive Cells	Cord Blood			ZAP Express
H0306	CD34 depleted Buffy Coat (Cord Blood)	CD34 Depleted Buffy Coat (Cord Blood)	Cord Blood			ZAP Express
H0309	Human Chronic Synovitis	Synovium, Chronic Synovitis/ Osteoarthritis	Synovium		disease	Uni-ZAP XR
H0316	HUMAN STOMACH	Human Stomach	Stomach			Uni-ZAP XR
H0318	HUMAN B CELL LYMPHOMA	Human B Cell Lymphoma	Lymph Node		disease	Uni-ZAP XR
H0321	HUMAN SCHWANOMA	Schwannoma	Nerve		disease	Uni-ZAP XR
H0327	human corpus colosum	Human Corpus Callosum	Brain			Uni-ZAP XR
H0328	human ovarian cancer	Ovarian Cancer	Ovary		disease	Uni-ZAP XR
H0329	Dermatofibrosarcoma Protuberance	Dermatofibrosarcoma Protuberans	Skin		disease	Uni-ZAP XR
H0331	Hepatocellular Tumor	Hepatocellular Tumor	Liver		disease	Lambda ZAP II
H0333	Hemangiopericytoma	Hemangiopericytoma	Blood vessel		disease	Lambda ZAP II
H0340	Corpus Callosum	Corpus Callosum-93052				Uni-ZAP XR
H0341	Bone Marrow Cell Line (RS4;11)	Bone Marrow Cell Line RS4;11	Bone Marrow	Cell Line		Uni-ZAP XR
H0343	stomach cancer (human)	Stomach Cancer - 5383A (human)			disease	Uni-ZAP XR
H0349	human adult liver cDNA library	Human Adult Liver	Liver			pCMVSPORT 1
H0351	Glioblastoma	Glioblastoma	Brain		disease	Uni-ZAP XR
H0352	wilm's tumor	Wilm's Tumor			disease	Uni-ZAP

	subtracted					
H0213	Human Pituitary, subtracted	Human Pituitary				Uni-ZAP XR
H0214	Raji cells, cyclohexamide treated, subtracted	Cyclohexamide Treated Cem, Jurkat, Raji, and Supt	Blood	Cell Line		pBluescript
H0216	Supt cells, cyclohexamide treated, subtracted	Cyclohexamide Treated Cem, Jurkat, Raji, and Supt	Blood	Cell Line		pBluescript
H0217	Supt cells, cyclohexamide treated, differentially expressed	Cyclohexamide Treated Cem, Jurkat, Raji, and Supt	Blood	Cell Line		pBluescript
H0221	Activated T-Cells, 4 hrs, differentially expressed	Activated T-Cells	Blood	Cell Line		Uni-ZAP XR
H0222	Activated T-Cells, 8 hrs, subtracted	Activated T-Cells	Blood	Cell Line		Uni-ZAP XR
H0229	Early Stage Human Brain, random primed	Early Stage Human Brain	Brain			Lambda ZAP II
H0230	Human Cardiomyopathy, diff exp	Human Cardiomyopathy	Heart		disease	Uni-ZAP XR
H0231	Human Colon, subtraction	Human Colon				pBluescript
H0234	human colon cancer, metastatic to liver, differentially expressed	Human Colon Cancer, metastaticized to liver	Liver			pBluescript
H0235	Human colon cancer, metaticized to liver, subtraction	Human Colon Cancer, metastaticized to liver	Liver			pBluescript
H0244	Human 8 Week Whole Embryo, subtracted	Human 8 Week Old Embryo	Embryo			Uni-ZAP XR
H0250	Human Activated Monocytes	Human Monocytes				Uni-ZAP XR
H0251	Human Chondrosarcoma	Human Chondrosarcoma	Cartilage		disease	Uni-ZAP XR
H0252	Human Osteosarcoma	Human Osteosarcoma	Bone		disease	Uni-ZAP XR
H0253	Human adult testis, large inserts	Human Adult Testis	Testis			Uni-ZAP XR
H0254	Breast Lymph node cDNA library	Breast Lymph Node	Lymph Node			Uni-ZAP XR
H0255	breast lymph node CDNA library	Breast Lymph Node	Lymph Node			Lambda ZAP II
H0257	HL-60, PMA 4H	HL-60 Cells, PMA stimulated 4H	Blood	Cell Line		Uni-ZAP XR
H0261	H. cerebellum, Enzyme subtracted	Human Cerebellum	Brain			Uni-ZAP XR
H0263	human colon cancer	Human Colon Cancer	Colon		disease	Lambda ZAP II
H0264	human tonsils	Human Tonsil	Tonsil			Uni-ZAP XR
H0265	Activated T-Cell (12hs)/Thiouridine labelledEco	T-Cells	Blood	Cell Line		Uni-ZAP XR
H0266	Human Microvascular Endothelial Cells, fract. A	HMEC	Vein	Cell Line		Lambda ZAP II
H0267	Human Microvascular Endothelial Cells, fract. B	HMEC	Vein	Cell Line		Lambda ZAP II
H0268	Human Umbilical Vein Endothelial Cells, fract.	HUVE Cells	Umbilical vein	Cell Line		Lambda ZAP II

	Human, subtracted	Week Old Embryo				XR
H0150	Human Epididymus	Epididymis	Testis			Uni-ZAP XR
H0151	Early Stage Human Liver	Human Fetal Liver	Liver			Uni-ZAP XR
H0154	Human Fibrosarcoma	Human Skin Fibrosarcoma	Skin		disease	Uni-ZAP XR
H0155	Human Thymus, subtracted	Human Thymus Tumor	Thymus			pBluescript
H0156	Human Adrenal Gland Tumor	Human Adrenal Gland Tumor	Adrenal Gland		disease	Uni-ZAP XR
H0158	Activated T-Cells, 4 hrs., ligation 2	Activated T-Cells	Blood	Cell Line		Uni-ZAP XR
H0159	Activated T-Cells, 8 hrs., ligation 2	Activated T-Cells	Blood	Cell Line		Uni-ZAP XR
H0163	Human Synovium	Human Synovium	Synovium			Uni-ZAP XR
H0166	Human Prostate Cancer, Stage B2 fraction	Human Prostate Cancer, stage B2	Prostate		disease	Uni-ZAP XR
H0169	Human Prostate Cancer, Stage C fraction	Human Prostate Cancer, stage C	Prostate		disease	Uni-ZAP XR
H0170	12 Week Old Early Stage Human	Twelve Week Old Early Stage Human	Embryo			Uni-ZAP XR
H0171	12 Week Old Early Stage Human, II	Twelve Week Old Early Stage Human	Embryo			Uni-ZAP XR
H0172	Human Fetal Brain, random primed	Human Fetal Brain	Brain			Lambda ZAP II
H0177	CAMA1Ee Cell Line	CAMA1Ee Cell Line	Breast	Cell Line		Uni-ZAP XR
H0178	Human Fetal Brain	Human Fetal Brain	Brain			Uni-ZAP XR
H0179	Human Neutrophil	Human Neutrophil	Blood	Cell Line		Uni-ZAP XR
H0181	Human Primary Breast Cancer	Human Primary Breast Cancer	Breast		disease	Uni-ZAP XR
H0182	Human Primary Breast Cancer	Human Primary Breast Cancer	Breast		disease	Uni-ZAP XR
H0184	Human Colon Cancer, metastasized to liver	Human Colon Cancer, metastasized to liver	Liver		disease	Lambda ZAP II
H0187	Resting T-Cell	T-Cells	Blood	Cell Line		Lambda ZAP II
H0188	Human Normal Breast	Human Normal Breast	Breast			Uni-ZAP XR
H0192	Cem Cells, cyclohexamide treated, subtra	Cyclohexamide Treated Cem, Jurkat, Raji, and Supt	Blood	Cell Line		Uni-ZAP XR
H0194	Human Cerebellum, subtracted	Human Cerebellum	Brain			pBluescript
H0196	Human Cardiomyopathy, subtracted	Human Cardiomyopathy	Heart			Uni-ZAP XR
H0204	Human Colon Cancer, subtracted	Human Colon Cancer	Colon			pBluescript
H0208	Early Stage Human Lung, subtracted	Human Fetal Lung	Lung			pBluescript
H0209	Human Cerebellum, differentially expressed	Human Cerebellum	Brain			Uni-ZAP XR
H0211	Human Prostate, differential expression	Human Prostate	Prostate			pBluescript
H0212	Human Prostate,	Human Prostate	Prostate			pBluescript

H0081	Human Fetal Epithelium (Skin)	Human Fetal Skin	Skin			Uni-ZAP XR
H0082	Human Fetal Muscle	Human Fetal Muscle	Sk Muscle			Uni-ZAP XR
H0083	HUMAN JURKAT MEMBRANE BOUND POLYSOMES	Jurkat Cells				Uni-ZAP XR
H0085	Human Colon	Human Colon				Lambda ZAP II
H0086	Human epithelioid sarcoma	Epithelioid Sarcoma, muscle	Sk Muscle		disease	Uni-ZAP XR
H0087	Human Thymus	Human Thymus				pBluescript
H0090	Human T-Cell Lymphoma	T-Cell Lymphoma	T-Cell		disease	Uni-ZAP XR
H0097	Human Adult Heart, subtracted	Human Adult Heart	Heart			pBluescript
H0098	Human Adult Liver, subtracted	Human Adult Liver	Liver			Uni-ZAP XR
H0099	Human Lung Cancer, subtracted	Human Lung Cancer	Lung			pBluescript
H0100	Human Whole Six Week Old Embryo	Human Whole Six Week Old Embryo	Embryo			Uni-ZAP XR
H0102	Human Whole 6 Week Old Embryo (II), subt	Human Whole Six Week Old Embryo	Embryo			pBluescript
H0105	Human Fetal Heart, subtracted	Human Fetal Heart	Heart			pBluescript
H0109	Human Macrophage, subtracted	Macrophage	Blood	Cell Line		pBluescript
H0112	Human Parathyroid Tumor, subtracted	Human Parathyroid Tumor	Parathyroid			pBluescript
H0116	Human Thymus Tumor, subtracted	Human Thymus Tumor	Thymus			pBluescript
H0119	Human Pediatric Kidney	Human Pediatric Kidney	Kidney			Uni-ZAP XR
H0122	Human Adult Skeletal Muscle	Human Skeletal Muscle	Sk Muscle			Uni-ZAP XR
H0123	Human Fetal Dura Mater	Human Fetal Dura Mater	Brain			Uni-ZAP XR
H0124	Human Rhabdomyosarcoma	Human Rhabdomyosarcoma	Sk Muscle		disease	Uni-ZAP XR
H0125	Cem cells cyclohexamide treated	Cyclohexamide Treated Cem, Jurkat, Raji, and Supt	Blood	Cell Line		Uni-ZAP XR
H0129	Jurkat cells, thiouridine activated, fract II	Jurkat Cells				Uni-ZAP XR
H0130	LNCAP untreated	LNCAP Cell Line	Prostate	Cell Line		Uni-ZAP XR
H0131	LNCAP + 0.3nM R1881	LNCAP Cell Line	Prostate	Cell Line		Uni-ZAP XR
H0132	LNCAP + 30nM R1881	LNCAP Cell Line	Prostate	Cell Line		Uni-ZAP XR
H0134	Raji Cells, cyclohexamide treated	Cyclohexamide Treated Cem, Jurkat, Raji, and Supt	Blood	Cell Line		Uni-ZAP XR
H0135	Human Synovial Sarcoma	Human Synovial Sarcoma	Synovium			Uni-ZAP XR
H0136	Supt Cells, cyclohexamide treated	Cyclohexamide Treated Cem, Jurkat, Raji, and Supt	Blood	Cell Line		Uni-ZAP XR
H0144	Nine Week Old Early Stage Human	9 Wk Old Early Stage Human	Embryo			Uni-ZAP XR
H0149	7 Week Old Early Stage	Human Whole 7	Embryo			Uni-ZAP

	Embryo	Embryo				XR
H0014	Human Gall Bladder	Human Gall Bladder	Gall Bladder			Uni-ZAP XR
H0015	Human Gall Bladder, fraction II	Human Gall Bladder	Gall Bladder			Uni-ZAP XR
H0019	Human Fetal Heart	Human Fetal Heart	Heart			pBluescript
H0020	Human Hippocampus	Human Hippocampus	Brain			Uni-ZAP XR
H0021	Human Infant Adrenal Gland	Human Infant Adrenal Gland	Adrenal gland			Uni-ZAP XR
H0024	Human Fetal Lung III	Human Fetal Lung	Lung			Uni-ZAP XR
H0030	Human Placenta					Uni-ZAP XR
H0031	Human Placenta	Human Placenta	Placenta			Uni-ZAP XR
H0032	Human Prostate	Human Prostate	Prostate			Uni-ZAP XR
H0036	Human Adult Small Intestine	Human Adult Small Intestine	Small Int.			Uni-ZAP XR
H0038	Human Testes	Human Testes	Testis			Uni-ZAP XR
H0039	Human Pancreas Tumor	Human Pancreas Tumor	Pancreas		disease	Uni-ZAP XR
H0040	Human Testes Tumor	Human Testes Tumor	Testis		disease	Uni-ZAP XR
H0041	Human Fetal Bone	Human Fetal Bone	Bone			Uni-ZAP XR
H0042	Human Adult Pulmonary	Human Adult Pulmonary	Lung			Uni-ZAP XR
H0046	Human Endometrial Tumor	Human Endometrial Tumor	Uterus		disease	Uni-ZAP XR
H0048	Human Pineal Gland	Human Pineal Gland				Uni-ZAP XR
H0050	Human Fetal Heart	Human Fetal Heart	Heart			Uni-ZAP XR
H0051	Human Hippocampus	Human Hippocampus	Brain			Uni-ZAP XR
H0052	Human Cerebellum	Human Cerebellum	Brain			Uni-ZAP XR
H0053	Human Adult Kidney	Human Adult Kidney	Kidney			Uni-ZAP XR
H0056	Human Umbilical Vein, Endo. remake	Human Umbilical Vein Endothelial Cells	Umbilical vein			Uni-ZAP XR
H0057	Human Fetal Spleen					Uni-ZAP XR
H0059	Human Uterine Cancer	Human Uterine Cancer	Uterus		disease	Lambda ZAP II
H0060	Human Macrophage	Human Macrophage	Blood	Cell Line		pBluescript
H0062	Human Thymus	Human Thymus	Thymus			Uni-ZAP XR
H0063	Human Thymus	Human Thymus	Thymus			Uni-ZAP XR
H0068	Human Skin Tumor	Human Skin Tumor	Skin		disease	Uni-ZAP XR
H0069	Human Activated T-Cells	Activated T-Cells	Blood	Cell Line		Uni-ZAP XR
H0071	Human Infant Adrenal Gland	Human Infant Adrenal Gland	Adrenal gland			Uni-ZAP XR
H0074	Human Platelets	Human Platelets	Blood	Cell Line		Uni-ZAP XR

AR286	Skeletal Muscle (42386)	Skeletal Muscle (42386)				
AR287	Skeletal Muscle (42461)	Skeletal Muscle (42461)				
AR288	Skeletal Muscle (91-01-A105)	Skeletal Muscle (91-01-A105)				
AR289	Skeletal Muscle (92-04-A059)	Skeletal Muscle (92-04-A059)				
AR290	Skeletal Muscle (96-08-A171)	Skeletal Muscle (96-08-A171)				
AR291	Skeletal Muscle (97-07-A190A)	Skeletal Muscle (97-07-A190A)				
AR292	Skeletal Muscle Diabetic (42352)	Skeletal Muscle Diabetic (42352)				
AR293	Skeletal Muscle Diabetic (42366)	Skeletal Muscle Diabetic (42366)				
AR294	Skeletal Muscle Diabetic (42395)	Skeletal Muscle Diabetic (42395)				
AR295	Skeletal Muscle Diabetic (42483)	Skeletal Muscle Diabetic (42483)				
AR296	Skeletal Muscle Diabetic (42491)	Skeletal Muscle Diabetic (42491)				
AR297	Skeletal Muscle Diabetic 42352	Skeletal Muscle Diabetic 42352				
AR298	Skeletal Muscle (42461)	Skeletal Muscle (42461)				
AR299	Small Intestine	Small Intestine				
AR300	Stomach	Stomach				
AR301	T-Cell + HDPBQ71.fc 1449 16hrs	T-Cell + HDPBQ71.fc 1449 16hrs				
AR302	T-Cell + HDPBQ71.fc 1449 6hrs	T-Cell + HDPBQ71.fc 1449 6hrs				
AR303	T-Cell + IL2 16hrs	T-Cell + IL2 16hrs				
AR304	T-Cell + IL2 6hrs	T-Cell + IL2 6hrs				
AR306	T-Cell Untreated 16hrs	T-Cell Untreated 16hrs				
AR307	T-Cell Untreated 6hrs	T-Cell Untreated 6hrs				
AR308	T-Cells 24 hours	T-Cells 24 hours				
AR309	T-Cells 24 hrs	T-Cells 24 hrs				
AR310	T-Cells 24 hrs.	T-Cells 24 hrs.				
AR311	T-Cells 24hrs	T-Cells 24hrs				
AR312	T-Cells 4 days	T-Cells 4 days				
AR313	Thymus	Thymus				
AR314	TRE	TRE				
AR315	TREC	TREC				
AR316	Virtual Mixture	Virtual Mixture				
H0002	Human Adult Heart	Human Adult Heart	Heart			Uni-ZAP XR
H0004	Human Adult Spleen	Human Adult Spleen	Spleen			Uni-ZAP XR
H0007	Human Cerebellum	Human Cerebellum	Brain			Uni-ZAP XR
H0008	Whole 6 Week Old Embryo					Uni-ZAP XR
H0009	Human Fetal Brain					Uni-ZAP XR
H0012	Human Fetal Kidney	Human Fetal Kidney	Kidney			Uni-ZAP XR
H0013	Human 8 Week Whole	Human 8 Week Old	Embryo			Uni-ZAP

AR248	Monocyte (CT)	Monocyte (CT)				
AR249	Monocyte (OCT)	Monocyte (OCT)				
AR250	Monocytes (CT)	Monocytes (CT)				
AR251	Monocytes (INFG 18 hr)	Monocytes (INFG 18 hr)				
AR252	Monocytes (INFG 18hr)	Monocytes (INFG 18hr)				
AR253	Monocytes (INFG 8-11)	Monocytes (INFG 8-11)				
AR254	Monocytes (O CT)	Monocytes (O CT)				
AR255	Muscle (91-01-A105)	Muscle (91-01-A105)				
AR256	Muscle (92-04-A059)	Muscle (92-04-A059)				
AR257	Muscle (97-11-A056d)	Muscle (97-11-A056d)				
AR258	Muscle (99-06-A210A)	Muscle (99-06-A210A)				
AR259	Muscle (99-07-A203B)	Muscle (99-07-A203B)				
AR260	Muscle (99-7-A203B)	Muscle (99-7-A203B)				
AR261	Muscle Diabetic (42352R)	Muscle Diabetic (42352R)				
AR262	Muscle Diabetic (42366)	Muscle Diabetic (42366)				
AR263	NK-19 Control	NK-19 Control				
AR264	NK-19 IL Treated 72hrs	NK-19 IL Treated 72hrs				
AR265	NK-19 UK Treated 72 hrs.	NK-19 UK Treated 72 hrs.				
AR266	Omentum Normal (94-08-B009)	Omentum Normal (94-08-B009)				
AR267	Omentum Normal (97-01-A039A)	Omentum Normal (97-01-A039A)				
AR268	Omentum Normal (97-04-A114C)	Omentum Normal (97-04-A114C)				
AR269	Omentum Normal (97-06-A117C)	Omentum Normal (97-06-A117C)				
AR270	Omentum Normal (97-09-B004C)	Omentum Normal (97-09-B004C)				
AR271	Ovarian Cancer (17717AID)	Ovarian Cancer (17717AID)				
AR272	Ovarian Cancer (9905C023RC)	Ovarian Cancer (9905C023RC)				
AR273	Ovarian Cancer (9905C032RC)	Ovarian Cancer (9905C032RC)				
AR274	Ovary (9508G045)	Ovary (9508G045)				
AR275	Ovary (9701G208)	Ovary (9701G208)				
AR276	Ovary 9806G005	Ovary 9806G005				
AR277	Pancreas	Pancreas				
AR278	Placebo	Placebo				
AR279	rIL2 Control	rIL2 Control				
AR280	RSS288L	RSS288L				
AR281	RSS288LC	RSS288LC				
AR282	Salivary Gland	Salivary Gland				
AR283	Skeletal Muscle	Skeletal Muscle				
AR284	Skeletal Muscle (91-01-A105)	Skeletal Muscle (91-01-A105)				
AR285	Skeletal Muscle (42180)	Skeletal Muscle (42180)				

AR210	Cultured Preadipocyte (blue)	Cultured Preadipocyte (blue)				
AR211	Cultured Preadipocyte (Red)	Cultured Preadipocyte (Red)				
AR212	Donor II B-Cells 24hrs	Donor II B-Cells 24hrs				
AR213	Donor II Resting B-Cells	Donor II Resting B-Cells				
AR214	H114EP12 10nM Insulin	H114EP12 10nM Insulin				
AR215	H114EP12 (10nM insulin)	H114EP12 (10nM insulin)				
AR216	H114EP12 (2.6ug/ul)	H114EP12 (2.6ug/ul)				
AR217	H114EP12 (3.6ug/ul)	H114EP12 (3.6ug/ul)				
AR218	HUVEC #1	HUVEC #1				
AR219	HUVEC #2	HUVEC #2				
AR221	L6 undiff.	L6 undiff.				
AR222	L6 Undifferentiated	L6 Undifferentiated				
AR223	L6P8 + 10nM Insulin	L6P8 + 10nM Insulin				
AR224	L6P8 + HS	L6P8 + HS				
AR225	L6P8 10nM Insulin	L6P8 10nM Insulin				
AR226	Liver (00-06-A007B)	Liver (00-06-A007B)				
AR227	Liver (96-02-A075)	Liver (96-02-A075)				
AR228	Liver (96-03-A144)	Liver (96-03-A144)				
AR229	Liver (96-04-A138)	Liver (96-04-A138)				
AR230	Liver (97-10-A074B)	Liver (97-10-A074B)				
AR231	Liver (98-09-A242A)	Liver (98-09-A242A)				
AR232	Liver Diabetic (1042)	Liver Diabetic (1042)				
AR233	Liver Diabetic (41616)	Liver Diabetic (41616)				
AR234	Liver Diabetic (41955)	Liver Diabetic (41955)				
AR235	Liver Diabetic (42352R)	Liver Diabetic (42352R)				
AR236	Liver Diabetic (42366)	Liver Diabetic (42366)				
AR237	Liver Diabetic (42483)	Liver Diabetic (42483)				
AR238	Liver Diabetic (42491)	Liver Diabetic (42491)				
AR239	Liver Diabetic (99-09-A281A)	Liver Diabetic (99-09-A281A)				
AR240	Lung	Lung				
AR241	Lung (27270)	Lung (27270)				
AR242	Lung (2727Q)	Lung (2727Q)				
AR243	Lung Cancer (4005116A1)	Lung Cancer (4005116A1)				
AR244	Lung Cancer (4005121A5)	Lung Cancer (4005121A5)				
AR245	Lung Cancer (4005121A5))	Lung Cancer (4005121A5))				
AR246	Lung Cancer (4005340A4)	Lung Cancer (4005340A4)				
AR247	Mammary Gland	Mammary Gland				

AR175	Adipose Diabetic (41661)	Adipose Diabetic (41661)				
AR176	Adipose Diabetic (41689)	Adipose Diabetic (41689)				
AR177	Adipose Diabetic (41706)	Adipose Diabetic (41706)				
AR178	Adipose Diabetic (42352)	Adipose Diabetic (42352)				
AR179	Adipose Diabetic (42366)	Adipose Diabetic (42366)				
AR180	Adipose Diabetic (42452)	Adipose Diabetic (42452)				
AR181	Adipose Diabetic (42491)	Adipose Diabetic (42491)				
AR182	Adipose Normal (41843)	Adipose Normal (41843)				
AR183	Adipose Normal (41893)	Adipose Normal (41893)				
AR184	Adipose Normal (42452)	Adipose Normal (42452)				
AR185	Adrenal Gland	Adrenal Gland				
AR186	Adrenal Gland + Whole Brain	Adrenal Gland + Whole Brain				
AR187	B7(1hr)+ (inverted)	B7(1hr)+ (inverted)				
AR188	Breast (18275A2B)	Breast (18275A2B)				
AR189	Breast (4004199)	Breast (4004199)				
AR190	Breast (4004399)	Breast (4004399)				
AR191	Breast (4004943B7)	Breast (4004943B7)				
AR192	Breast (4005570B1)	Breast (4005570B1)				
AR193	Breast Cancer (4004127A30)	Breast Cancer (4004127A30)				
AR194	Breast Cancer (400443A21)	Breast Cancer (400443A21)				
AR195	Breast Cancer (4004643A2)	Breast Cancer (4004643A2)				
AR196	Breast Cancer (4004710A7)	Breast Cancer (4004710A7)				
AR197	Breast Cancer (4004943A21)	Breast Cancer (4004943A21)				
AR198	Breast Cancer (400553A2)	Breast Cancer (400553A2)				
AR199	Breast Cancer (9805C046R)	Breast Cancer (9805C046R)				
AR200	Breast Cancer (9806C012R)	Breast Cancer (9806C012R)				
AR201	Breast Cancer (ODQ 45913)	Breast Cancer (ODQ 45913)				
AR202	Breast Cancer (ODQ45913)	Breast Cancer (ODQ45913)				
AR203	Breast Cancer (ODQ4591B)	Breast Cancer (ODQ4591B)				
AR204	Colon Cancer (15663)	Colon Cancer (15663)				
AR205	Colon Cancer (4005144A4)	Colon Cancer (4005144A4)				
AR206	Colon Cancer (4005413A4)	Colon Cancer (4005413A4)				
AR207	Colon Cancer (4005570B1)	Colon Cancer (4005570B1)				
AR208	Control RNA #1	Control RNA #1				
AR209	Control RNA #2	Control RNA #2				

AR096	Spleen	Spleen				
AR097	Thymus T cells activated	Thymus T cells activated				
AR098	Thymus T cells resting	Thymus T cells resting				
AR099	Tonsil	Tonsil				
AR100	Tonsil germinal center centroblast	Tonsil germinal center centroblast				
AR101	Tonsil germinal center B cell	Tonsil germinal center B cell				
AR102	Tonsil lymph node	Tonsil lymph node				
AR103	Tonsil memory B cell	Tonsil memory B cell				
AR104	Whole Brain	Whole Brain				
AR105	Xenograft ES-2	Xenograft ES-2				
AR106	Xenograft SW626	Xenograft SW626				
AR119	001: IL-2	001: IL-2				
AR120	001: IL-2.1	001: IL-2.1				
AR121	001: IL-2_b	001: IL-2_b				
AR124	002 : Monocytes untreated (1hr)	002 : Monocytes untreated (1hr)				
AR125	002 : Monocytes untreated (5hrs)	002 : Monocytes untreated (5hrs)				
AR126	002: Control.1C	002: Control.1C				
AR127	002: IL2.1C	002: IL2.1C				
AR130	003 : Placebo-treated Rat Lacrimal Gland	003 : Placebo-treated Rat Lacrimal Gland				
AR131	003 : Placebo-treated Rat Submandibular Gland	003 : Placebo-treated Rat Submandibular Gland				
AR135	004 : Monocytes untreated (5hrs)	004 : Monocytes untreated (5hrs)				
AR136	004 : Monocytes untreated 1hr	004 : Monocytes untreated 1hr				
AR139	005: Placebo (48hrs)	005: Placebo (48hrs)				
AR140	006: pC4 (24hrs)	006: pC4 (24hrs)				
AR141	006: pC4 (48hrs)	006: pC4 (48hrs)				
AR152	007: PHA(1hr)	007: PHA(1hr)				
AR153	007: PHA(6HRS)	007: PHA(6HRS)				
AR154	007: PMA(6hrs)	007: PMA(6hrs)				
AR155	008: 1449_#2	008: 1449_#2				
AR161	01: A - max 24	01: A - max 24				
AR162	01: A - max 26	01: A - max 26				
AR163	01: A - max 30	01: A - max 30				
AR164	01: B - max 24	01: B - max 24				
AR165	01: B - max 26	01: B - max 26				
AR166	01: B - max 30	01: B - max 30				
AR167	1449 Sample	1449 Sample				
AR168	3T3P10 1.0uM insulin	3T3P10 1.0uM insulin				
AR169	3T3P10 10nM Insulin	3T3P10 10nM Insulin				
AR170	3T3P10 10uM insulin	3T3P10 10uM insulin				
AR171	3T3P10 No Insulin	3T3P10 No Insulin				
AR172	3T3P4	3T3P4				
AR173	Adipose (41892)	Adipose (41892)				
AR174	Adipose Diabetic (41611)	Adipose Diabetic (41611)				

	lymphocytic leukaemia	chronic lymphocytic leukaemia				
AR064	Lymphocytes diffuse large B cell lymphoma	Lymphocytes diffuse large B cell lymphoma				
AR065	Lymphocytes follicular lymphoma	Lymphocytes follicular lymphoma				
AR066	normal breast	normal breast				
AR067	Normal Ovarian (4004901)	Normal Ovarian (4004901)				
AR068	Normal Ovary 9508G045	Normal Ovary 9508G045				
AR069	Normal Ovary 9701G208	Normal Ovary 9701G208				
AR070	Normal Ovary 9806G005	Normal Ovary 9806G005				
AR071	Ovarian Cancer	Ovarian Cancer				
AR072	Ovarian Cancer (9702G001)	Ovarian Cancer (9702G001)				
AR073	Ovarian Cancer (9707G029)	Ovarian Cancer (9707G029)				
AR074	Ovarian Cancer (9804G011)	Ovarian Cancer (9804G011)				
AR075	Ovarian Cancer (9806G019)	Ovarian Cancer (9806G019)				
AR076	Ovarian Cancer (9807G017)	Ovarian Cancer (9807G017)				
AR077	Ovarian Cancer (9809G001)	Ovarian Cancer (9809G001)				
AR078	ovarian cancer 15799	ovarian cancer 15799				
AR079	Ovarian Cancer 17717AID	Ovarian Cancer 17717AID				
AR080	Ovarian Cancer 4004664B1	Ovarian Cancer 4004664B1				
AR081	Ovarian Cancer 4005315A1	Ovarian Cancer 4005315A1				
AR082	ovarian cancer 94127303	ovarian cancer 94127303				
AR083	Ovarian Cancer 96069304	Ovarian Cancer 96069304				
AR084	Ovarian Cancer 9707G029	Ovarian Cancer 9707G029				
AR085	Ovarian Cancer 9807G045	Ovarian Cancer 9807G045				
AR086	ovarian cancer 9809G001	ovarian cancer 9809G001				
AR087	Ovarian Cancer 9905C032RC	Ovarian Cancer 9905C032RC				
AR088	Ovarian cancer 9907 C00 3rd	Ovarian cancer 9907 C00 3rd				
AR089	Prostate	Prostate				
AR090	Prostate (clonotech)	Prostate (clonotech)				
AR091	prostate cancer	prostate cancer				
AR092	prostate cancer #15176	prostate cancer #15176				
AR093	prostate cancer #15509	prostate cancer #15509				
AR094	prostate cancer #15673	prostate cancer #15673				
AR095	Small Intestine (Clontech)	Small Intestine (Clontech)				

TABLE 4

Code	Description	Tissue	Organ	Cell Line	Disease	Vector
AR022	a_Heart	a_Heart				
AR023	a_Liver	a_Liver				
AR024	a_mammary gland	a_mammary gland				
AR025	a_Prostate	a_Prostate				
AR026	a_small intestine	a_small intestine				
AR027	a_Stomach	a_Stomach				
AR028	Blood B cells	Blood B cells				
AR029	Blood B cells activated	Blood B cells activated				
AR030	Blood B cells resting	Blood B cells resting				
AR031	Blood T cells activated	Blood T cells activated				
AR032	Blood T cells resting	Blood T cells resting				
AR033	brain	brain				
AR034	breast	breast				
AR035	breast cancer	breast cancer				
AR036	Cell Line CAO V3	Cell Line CAO V3				
AR037	cell line PA-1	cell line PA-1				
AR038	cell line transformed	cell line transformed				
AR039	colon	colon				
AR040	colon (9808co65R)	colon (9808co65R)				
AR041	colon (9809co15)	colon (9809co15)				
AR042	colon cancer	colon cancer				
AR043	colon cancer (9808co64R)	colon cancer (9808co64R)				
AR044	colon cancer 9809co14	colon cancer 9809co14				
AR045	corn clone 5	corn clone 5				
AR046	corn clone 6	corn clone 6				
AR047	corn clone2	corn clone2				
AR048	corn clone3	corn clone3				
AR049	Corn Clone4	Corn Clone4				
AR050	Donor II B Cells 24hrs	Donor II B Cells 24hrs				
AR051	Donor II B Cells 72hrs	Donor II B Cells 72hrs				
AR052	Donor II B-Cells 24 hrs.	Donor II B-Cells 24 hrs.				
AR053	Donor II B-Cells 72hrs	Donor II B-Cells 72hrs				
AR054	Donor II Resting B Cells	Donor II Resting B Cells				
AR055	Heart	Heart				
AR056	Human Lung (clonotech)	Human Lung (clonotech)				
AR057	Human Mammary (clonotech)	Human Mammary (clonotech)				
AR058	Human Thymus (clonotech)	Human Thymus (clonotech)				
AR059	Jurkat (unstimulated)	Jurkat (unstimulated)				
AR060	Kidney	Kidney				
AR061	Liver	Liver				
AR062	Liver (Clontech)	Liver (Clontech)				
AR063	Lymphocytes chronic	Lymphocytes				

						BF996488, BF995754, BF086199, BF085946, AX013771, AL079279, AF206329, I76197, AK026038, and AF242376.
HTGDN55	156	1106269	1 - 641	15 - 655		AC003029.

HLHCR16	155	1106227	1 - 3294	15 - 3308	<p> AV720878, AV718844, AW973482, AV699447, AW958992, AW958993, AV723927, D81026, AW959136, AW964488, AW962082, AW949656, AW949642, AW959202, AA305578, C15076, AW966385, AV720151, D81030, AW973473, AW966378, D51060, AW966388, AW949500, AA514186, D80212, AW959469, AW966386, D80193, D59927, AW966368, AV720616, D80196, AW966331, D80188, AW966398, AW966397, AV699866, D80219, AW949658, AV719913, D59889, AW973330, AV700229, AW965158, AW966043, AV701839, AV718530, AW962245, D80366, AW960532, D80248, D80045, AW952852, D80522, D57483, AW177440, D80439, AW973447, AW966389, AV655880, AV720812, AV720533, AW964737, AW960454, AW966032, AW959582, D80247, AW966379, AW752082, AF192403, AF192402, A84916, AX047063, A62300, AX033851, AX027925, A62298, AR070327, AR087649, AJ132110, AB028859, A82595, AF058696, AR018138, AR008278, AR060385, AX047064, Y12724, AB002449, Y17188, AX021518, A30438, I50126, I50132, I50128, I50133, X67155, D26022, AX035434, A25909, A94995, AX028130, AX020191, A67220, D89785, A78862, D34614, AX020190, AJ302649, AR016514, AR016808, AR066488, AR060138, A26615, AR052274, A45456, AX047062, D88547, AR008443, Y09669, AR074139, A43192, A43190, AR038669, AR066487, AR074136, I14842, AR054175, X82626, Y17187, AR025207, AR008277, AR008281, D50010, A63261, AR077702, AR008408, AR091537, AR062872, A70867, AR093385, U46128, AR016691, AR016690, I18367, AR074545, AB012117, AR060133, I79511, A64136, A68321, A85396, AR074141, D88507, AR066482, A44171, AR088705, X68127, AX042372, T66450, and AW152154. </p> <p> AA402528, AI379350, AA716107, AI123557, BG012978, BF996491, BG012967, AW983663, AI127175, BE810281, AA234106, AA234698, BG010666, AI039768, BG012979, AI580137, N77999, AA424560, AA419490, AI334141, R71349, AI224976, AI417798, AI080508, BG012946, N58410, AI818475, N91089, BF986530, AA424657, AA399612, AI144265, AA399137, AI498363, BG013866, AA410986, AA235306, AA399148, BG010096, AW292497, BG012565, AI249102, W86869, AA115407, H81257, BG012976, BE817685, AI077499, H00194, H45499, BG012578, BG013065, BF998792, R01206, BF997284, AW104245, BG014612, BG009928, BG012573, AA234880, F05822, BF369785, AA852176, R82875, R71350, H81245, AA298421, R22096, AA367417, H81585, T49265, H81591, AI985171, AA115408, R64037, T54283, AI492930, BG014506, AI492932, R82876, R34138, R81695, R33935, R25630, BF992066, H81595, R81696, R32025, T49264, R95688, AW970150, AW970146, BG006854, R01319, BF993436, Z21579, R22040, R33043, Z38741, BF996770, BF085179, AA514393, T97662, BE817625, T34993, BG013537, BE930065, AA705441, BF107166, D79105, R32078, BF843592, BG012975, BE817684, BE817611, BF997299, H45500, AA707178, R30943, H81576, H81250, AI032721, BF996793, BF990036, </p>
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HDPMQ42	152	1106192	1 - 1312	15 - 1326	AI949910, AV716802, AW242656, AV716725, AI400631, AI338670, AA418441, AA446401, AA677149, AV711676, AI333747, C01555, AA380065, AA700481, AF067800, AI133532, AF109146, AF200738, and AC006511.
HHFHQ39	153	1106209	1 - 3613	15 - 3627	AI492875, AI796676, AI749838, AA918144, AI814590, AI923531, AW188476, AA524255, BF513992, AA847476, AA483519, AI720725, AI748873, AI150879, AI279072, AW190984, AI284510, AA775055, AW612904, AI492104, AI141231, AA613554, AW662148, AI917655, AW769047, AA662104, AA565985, T89953, AW581798, AI623640, AA377113, AI473980, AA295763, BE934311, AW769091, BF064110, AW974745, BF063227, AA295769, AA295332, AA662070, AA075641, BF849553, AA075642, BE829414, AW593671, AA295795, BF760645, AW612888, BE829378, AA309786, BE932867, AI626076, AW769569, AI796678, AI815239, AI045413, AW366372, AW149876, AI559752, AW079432, AI872423, BF344583, AI285417, AI567971, AL121496, AV738825, AV736475, AV737685, AV735391, AV743177, AV740451, AV735682, AV736885, AI357672, AV744590, AI927233, AV735798, AV743713, AV739235, AV745038, AI285439, AW151136, AI567582, AI539771, AI867068, AI952433, BE897632, AI432644, AI537677, AI494201, AI690813, BF812963, AI890907, AI804505, AI500659, BE883591, AI866465, AI815232, AI801325, AI866691, AI500523, BF812438, AI538850, BE885490, AI887775, AI582932, AI590043, AI923989, AI284517, AI500706, AI445237, AI491776, BF811804, AI289791, AW151138, AI926593, AI889189, AI521560, AI500662, AI284509, AI539800, AI582912, AW172723, AI889168, AI538885, AI440263, BG029399, AI866573, AI633493, AI434256, AF159456, A90822, AJ243224, AJ243212, AF105035, AJ243211, A90827, AF043112, AB005909, U37438, I72544, U32681, A90826, S78981, AB020851, AB020850, AB020833, AB020836, AB020830, AB020844, AB020839, AB020821, AB020827, AB020824, AB020818, AC005522, AB020848, AB020843, AB020849, AB020842, AF079224, AB020847, AK027164, X54971, AI050280, AF254119, I48978, AL110159, AL133049, AL133070, AF113690, AL050277, AK026603, AL389935, AK000603, AF080622, U86379, AL117443, Y17327, AF137370, D16301, AX040958, M85165, AX040974, U30290, AL390154, AF130092, AL137459, AL137478, AR022283, I89947,

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H7TSA02	147	1105584	1 - 692	15 - 706	AA077171, BF946981, BF314839, AR028901, AR028893, AR028894, AR028902, AC006466, AR028891, L16506, D17516, D82935, AR028892, AR028900, AR028895, L16680, AR028903, Z23274, AR028899, D14909, AR028888, Z23282, D16465, D17290, Z23272, Z23275, AR028890, Z23279, AR028898, D14908, Z23273, AR028889, and AR028897.
H7TSA19	148	1105585	1 - 497	15 - 511	BF946981, AA077171, D17516, AR028892, AR028900, AR028894, AR028893, AR028902, AR028901, AR028895, AR028903, AC006466, AR028890, Z23279, AR028898, D14908, AR028889, D82935, AR028897, L16680, L16506, AR028891, Z23275, Z23274, Z23272, Z23282, AR028899, D14909, D16465, Z23273, AR028888, and D17290.
HMWCU6 3	149	1105747	1 - 606	15 - 620	BF103655, AW408031, BE247532, AA380908, AA029932, BE929465, AW956578, AI718659, AI651912, BG232105, BE049585, AA749217, AI984467, AI633684, AA743043, AA748305, BF513581, BF895003, BF895254, AF285089, AC007068, AF133299, AF285087, AF285088, and AC010186.
HWADF26	150	1105857	1 - 1552	15 - 1566	AA426628, BG168646, BG117025, BG168645, BG253641, BG258222, BG167818, BG025417, BG164284, BG612896, BG252223, AW971745, BE910738, BF345060, BE621893, BE621811, BF984267, BG123011, BF798053, BF984992, BG031014, BE621810, BG117030, BE613349, BE612945, BE622828, BE613070, AA424848, BG117026, BE620742, AA173389, BE613136, BE622931, BE621911, BE614378, BE614307, BG168644, BE614237, BE622925, BE622868, BE618811, BE614451, BE614230, BG253645, T60164, BE613101,

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HKACI03	145	1104932	1 - 1883	15 - 1897	<p>BF909313, BE259856, BF316313, BF892536, BE747113, BF847845, BE273557, BG034698, BE302770, AA534193, AI529248, BE707826, BE533851, AL519618, BF846528, AW984086, BE938073, AI830367, BE727051, AI360265, BE392753, BE280593, AC005594, and AC005783.</p> <p>BE867597, BE563419, BE907794, BE792267, BE563484, BF916758, BE744931, BE743617, BE549271, BE250039, BE743983, BE740414, BF916759, BF970916, BE974183, BE541095, BE728270, BF196344, AI241578, BE746660, BE698579, BF037459, BG109047, BE698577, BE272401, BE698578, BF818581, BE698576, BE698574, BF916525, BE305089, BE709697, BF816476, BE866320, BF132539, BE568812, AI955238, AW804494, BE871927, AI312113, BE709749, BE871269, BF914938, BE927878, AI979207, AW338261, BE562400, BF770534, BF770846, AI925871, BF676007, BE561772, BE561159, AW009598, AA741250, AI333843, AI590808, BE350662, BE396947, T86963, AW190875, AW952036, N20044, BE562032, AA244018, AA101562, AA149993, BE562362, AW192453, BF820875, AA152150, BF833534, AW583700, N28398, BF677728, BF676772, BF818090, AA483522, BF846233, BG123012, BE162549, BE709576, T40695, BE866069, R72982, BG013078, BE268930, AA224590, AI093487, AA244075, AI263270, AA101561, BF771639, BF818208, T87045, AI193265, BF834943, R01692, AI469580, AW999404, BF871808, AI632044, W76302, T84017, BF836755, AW510529, BF871973, R28222, R02633, AI002582, BF818279, AA101520, BF803095, BF839692, AI864718, T39606, AA304161, BF813448, BF033572, BF868424, BE241748, BF813934, AA702259, AI608881, R27969, AW994103, T39607, BF811982, T84016, C75490, BG010167, C75621, BF873312, BF871813, AA861308, BE871109, AA890390, BF747862, AA486163, AW630757, BF825235, AA513370, AA404390, AA486100, BF747135, AW752957, BF832600, AA641435, AA227408, BF807059, AI687737, T73746, AA215609, AA642887, BF849699, AA101519, AI799264, C17760, H57108, AA768079, BF848642, BF133227, AI304788, BF825377, AW936241, AW996633, AI935684, AW339937, BF353664, BE927881, AF191495, AF207907, AF111713, AF154005, AK026665, AF172398, A91699, A91701, AF111714, AF207550, AC008498, AC008784, AL118520, AC004913, and AC026425.</p> <p>BF972832, BF968664, BF337801, BF038410, BF338126, BF688743, BF688560, BF339410, BF726248, BF338788, BF528351, BE615509, AI800218, AI346281, AW748162, BG222425, BF036565, AI983710, BF689301, AW026429, AI766300, BF805547, BG249715, AI583539, AW264048, AI911743, BG060162, BG117232, AW387854, AI924093, AW470078, AI264211, AI312053, AI201482, AI955668, BE615228, BE677699, AW058431, BE045835, AW196781, BE221691, AI022218, BF983475, AI565141, BE737528, AI689249, BF689198, AI276048, AI566956, BE736613, AI920904, BG056452, AW193090, AI623560, BE018907, AW664907, BF570443, AW615392, BE931661, BF588543, BG166108, AA922838,</p>
HDTEI03	146	1104936	1 - 2630	15 - 2644	

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HETAQ37	142	1103950	1 - 1946	15 - 1960		<p>AW956492, BF352284, BE709448, W29010, AV751953, AI829559, BE179448, AI571060, AI083491, AA905071, AW118125, AJ049799, W22553, N90902, W27632, AI273588, AI890622, AW195777, W27896, W22119, BF844755, AW269932, AA040604, W23268, W38638, W37154, AA904910, W27944, R55894, W27681, AW607334, AA337059, AW367713, AW607165, AW966017, BF351195, AA298658, R55809, AI376671, C02576, BE163743, AA364393, AV746728, AI393483, W23093, BE708746, W28670, W27851, BF844715, H83295, BF437730, N92239, H83294, AI194027, W27371, D81988, BE163611, C14616, C14877, BE163731, AA040705, AA897696, AV747434, BF850779, AA017680, AW974810, AF122922, and AF122923.</p>
HCENW09	143	1104186	1 - 731	15 - 745		<p>BG115463, BF795158, BE389127, BE390471, A1768326, AI333117, AA018361, BE388347, AA324901, BE305115, BE249961, BE842897, BE910034, AW674496, F07835, AW577798, AL117482, AR077242, AL360256, and Z61430.</p>
HAMFW0 5	144	1104784	1 - 2752	15 - 2766		<p>BE794483, BG178503, BG108176, BG167854, BF203783, AW374284, BE407965, BE296162, BF203610, AL519617, AL529247, BF897130, BF897134, BE813309, BF243324, BF683248, BE931606, BE707485, BF570610, BE785669, AW410587, BF930533, BF990414,</p>

	AI142999, AA081290, N93459, AA506965, AA401860, W25198, AI311111, R60726, R60727, AI351724, BF679941, AW614976, BF091172, AA604134, T63960, BE840632, AI351726, AW009121, AA081115, AA081697, AA411256, AW129067, AA578520, AA612772, AI824391, AA470674, H66783, AW953852, AI583100, N34727, AW160746, AA832062, R36715, AW770335, R90863, R84524, BF924179, H12158, AV738340, AV702770, AA315553, WW246146, AV742732, AV704180, AA370468, AV727978, AV728309, AW958796, AV728874, AV701643, AV706910, AA832305, AV704660, AV706223, AV703542, AV707329, AV706162, AV728436, AV702322, AV706882, AV726502, AV705684, AV727314, AV706851, AV706165, AV727954, AW082570, AV646347, AV727238, AV706147, AV660516, AW960067, AV705550, AV652528, AW955995, AV728404, AV709549, AV701728, AV661286, AV727583, AV725001, AV655096, AV709356, AV725255, AV707151, AV658784, AV725380, AV702164, AW955088, AW952301, AV706741, AV705518, AV707024, AV704144, AV703012, AV728715, AV742050, AV726392, AV727618, AV707830, AV703436, AV707639, AV658362, AW955629, AV703591, AV729242, AW955019, AV704307, AV725709, AV725386, AV702984, AV705416, AV701880, AV725577, AW959858, AW965148, AV725321, AV726590, AV703266, AW952013, AV709580, AV706925, AV727337, AW963446, AW956292, AV708520, AV729255, AV729597, AW958801, AV708872, AV705445, AV704756, AW957279, AV725393, AV727822, AV704857, AV725618, AV708786, AV725582, AV704401, AV653353, AW951526, AV688061, AV704626, AV651897, AV704558, AW950276, AV704568, AV702671, AV707589, AV729473, AV725152, AV658797, AW959986, AV725441, AV725031, AV702738, AV728289, AV704879, AV704077, AW949777, AW952497, AV709939, AV704712, AV729424, AV652001, AV704358, AV725930, AV728743, AV726337, AV731070, AV703762, AV728312, AV703035, AV725991, AV709025, AV706992, AV701874, AV726194, AV658436, AV726628, AV707658, AW967329, AV706683, AV661838, AV705538, AV703980, AV727047, AV728355, AV647654, AW960579, AV701626, AV708083, AL050275, AX047345, AX020190, AX047063, AR062871, AX047062, AX026824, AR070327, AX026823, AR062872, AR062873, A20702, AR017907, A20700, A43189, A43188, AR088705, AR038855, AX020191, AX047064, A62298, I19525, A84776, A84773, A84775, A84774, AR067731, AR067732, AS8522, A91750, AX035462, A62300, A25909, AX009487, AX001322, AX006816, AR091537, A85395, AR093385, A85476, Y09813, AR091393, Y17188, AR079804, A84916, U94592, AJ244005, Y16359, AX028305, AJ244004, AX001082, AX021518, AX006825, AX006822, AR037157, AX008555, AJ244003, AX006826, D50010, AX033851, AX027925, D78345, AR008429, I15717, AR054109, A98420, A98423, A98432, A98436, A98417, A98427, I15718, P0902424
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HDPSJ02	140	1103186	1 - 1616	15 - 1630	<p> BF898617, AA366176, AK024462, AK024459, AL109804, and AP002898. </p>
HE6FT04	141	1103911	1 - 2137	15 - 2151	<p> BF342508, BE745079, BF316647, AW954022, BF220093, BF219864, AI684112, BF182978, BG108443, BE879671, BE840525, AW957217, BE840530, AI148569, AI128199, AI041807, BF027688, BF915566, BF914452, AI991034, AI862148, AA588312, AA424398, AI333529, AI672251, AI798586, AI095534, BF826436, AI475525, AI039685, T52017, AI055912, R51437, AI598282, AW071787, AW028194, AA578538, AI140222, AI300146, AA554343, AW268634, AA340540, AA411182, AI076726, C04045, AW088744, AA757547, AV691484, AI798454, AW473352, AW249610, AV647523, AW475065, AA364829, W22554, H66782, </p>

HETJF84	138	1102857	1 - 2780	15 - 2794	<p>A92667, A92665, AR028791, AR028793, AX032758, AR018924, AR018923, A48774, A48775, AR000006, AR015960, AR015961, AR000007, AR051652, A91752, AR091393, AX008865, AX008867, AX008868, A92081, A92080, A92077, A92078, A92079, AX030369, AX030368, AR069417, AR069413, AX033488, AX033489, AX033490, AX033474, AX033486, AX033487, A91751, AX036660, AX036661, AR068508, AR068510, AR068509, AR091571, AR085090, AR003585, I58322, I58323, A85308, AX044426, AX044425, A91754, AX026821, A60213, AX046332, AR062871, A44171, AX027908, AX027904, AX027906, AR068550, A23373, AR068551, A49700, A60207, A60208, I58669, A58521, A63067, A51047, A63064, A63072, AR031375, AR068507, AR068506, AX001323, AX001324, A29109, A32111, AX004550, AR031374, I07209, I07249, A63954, AX030435, AX026820, AR019098, AR019097, AR020199, AR020200, AR001287, AR020198, AR020197, AR029418, AR067734, AR067731, AR067732, AR029417, AR051651, AX030436, I89986, AX024906, AR085093, AR051957, Y14971, I09121, Y17793, AR066494, AX046357, A93444, A46342, A46343, AB026436, AR055065, AR071207, AF064854, AF019249, AJ251859, AX040911, AR060234, A30600, AR032878, AR055066, A81671, AR054110, AJ279014, and AR069079.</p> <p>AV647669, BF694934, BF350634, AV647418, BF754372, T86312, BE142547, AA332746, AW754392, T93939, BE142562, and AL133553.</p>
HE2DL46	139	1102938	1 - 3014	15 - 3028	<p>BF220057, BE267201, BF309481, AW247289, BF220329, BE794949, BF832561, BF967189, AI808090, AW250783, AV728432, AV725167, BF222625, N47945, BF967916, BE267395, AA861657, AV723651, N77555, AA084165, AI038716, AW246922, AV732142, AI040877, D60694, AI023144, AA084138, AA694591, AL120454, BF205874, AW023630, AI984284, AA199586, N49154, AW975618, AV724520, AW949645, D59619, D80210, D80240, AV718844, AV718692, AW966531, D80219, AV718489, AW949656, AW949631, AW966053, AW978634, D58283, AV699550, AV742001, AW949642, D80212, AV720211, AW949643, D80195, AV723927, AW959628, D81030, D59927, D80391, AV719822, D51423, D51799, AV719324, AV719783, D80253, AV718800, AV720203, D80022, D80227, AV720464, AV718770, AW966062, AW959570, AV719188, AW949629, D80043, AW949632, AV720731, AV699447, D80193, D80196, AV722801, AW949641, AV719557, AV699927, D80188, AW975621, C14429, AW949646, AW949658, AW949657, D80038, AW949653, AV719468, AW965158, AW973307, AW973447, D80166, D80366, AW949618, AW949633, AV720812, AV721386, AW949655, D59889, AV720028, AW966013, AV723097, AW966534, AW960553, D80045, AV700889, AV718440, D59275, AW965177, D59859, AW966050, AW959582, AW966041, AW966054, AW949654, AW978661, AW973541, D80269, AV720791, AW959597, T03269, AV720654, AW959202, C14331, D80378, D59502, D50995, C75259, AV699746, AW960465, AW964488, AV701335,</p>

					AI460232, AI276126, AI128077, AA326017, BF345886, AA367412, AA126901, AA351643, BE789481, AW802995, BF755875, AL035681, and AL035681.
HRDFA94	134	1100389	1 - 1229	15 - 1243	AP002898, and AL109804.
HSIFQ16	135	1100483	1 - 2522	15 - 2536	AL529223, AI651405, AI660075, AI654857, AI431898, AI672616, AI624730, AW401851, AI624728, BE798665, BF594215, R46681, H26800, AI491709, AI570936, AA768926, AA650038, BF895967, AA806378, AW879618, AI623325, BF091262, AA310232, AI917017, BE646092, BG009720, AI350516, BE244625, BF594777, AW236839, BE621379, AW176705, BF795359, BE294675, BF797367, BG012781, AW016207, BE892810, AI992313, BG150450, AI631550, BF851617, AW197267, AL529222, N24032, BF375280, H14527, W07754, BG009905, AA948504, BF747812, AI825439, BE297459, AW182813, AA720646, BF741419, BF848728, AA947309, BG005101, AW575034, BF878062, BF739332, AF053356, and AR016407.
HWLDY64	136	1100528	1 - 1164	15 - 1178	BF334851, H68634, AI040942, H68655, H75859, AC002453, AC002453, and AC002453.
HELB20	137	1100919	1 - 2770	15 - 2784	AF150174, BF511602, AI417513, AI698235, AV755831, AV737500, AW592241, AV742237, BE841227, AW971745, AV681500, AV681492, AV645393, AV645392, AV645390, AV645385, AV681509, AV681529, AV681512, AV681526, AV645336, AV645317, AV681507, AV681465, AV681525, AV681495, AV681505, AV681502, AV681504, AV681486, AV681483, AV681464, AV681462, AV681487, AV681506, AV681523, AV681519, AV681471, AV681497, AV645358, AV681513, AV645401, AV645332, AV645383, AV645377, AV645379, AV645389, AV645369, AV645380, AV645321, AV681527, AV681501, AV645400, AV645404, AV645353, AV681458, AV645334, AV681510, AV681528, AV681474, AV681514, AV645339, AV681477, AV681491, AV681472, AV681475, AV681488, BE613349, BE612945, BE622828, BE613070, AV738011, BE614237, BE614307, BE613136, BE622931, BE621911, BE614230, BE622925, BE618811, BE622868, BE620742, AV681461, BE614451, AV645350, BE170947, AV645344, BE620756, AV681531, R56970, BF243224, AV755561, AV681468, AW972092, AW972093, AW861944, AW968355, AW968356, AW858525, AW968729, AW972091, AW877209, BF214634, AW971740, AI431351, AA471187, AW972090, AW969229, AW804686, AW858455, Z99396, AI432649, AL119324, AW604723, AI432644, AI623302, AI431337, AW858526, AI432674, AI432651, AI431241, AR080280, I25027, AR054109, I44515, I26928, I26930, I26927, I25041, AR069374, AR069375, I44516, AR091518, AX009487, AR093384, I85513, AR009152, AR009151, AR027099, AR093392, AR093383, AR035224, I05393, A10617, AR028792, A01324, A01323, A94046, A94054, AR064321, AR034783, AX030966, I63120, AR067733, AX009486, AX029455, AR064322, AR064323, AR064320, A94048, A94061, AX027785, AR038307, AR038321, I05430, A83642, A83643, AX035462, AX003194, A70359, A32110, AR083151, A49045, AR019094, A92666, A92668, I49890,

HARNY39	128	1097714	1 - 1580	15 - 1594	AR067734, AR067731, AR067732, AR029417, AX026820, AR019097, I89986, AX024906, AR085093, AR051651, AR051957, I09121, Y14971, AR026436, A46342, A46343, A93444, AR055065, AX030435, AJ251859, AX046357, AR060234, AX040911, AJ279014, A30600, AR055066, AR054110, AR032878, A81671, AR069079, AR066494, and AR043113.
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HIPDH20	129	1099283	1 - 1076	15 - 1090	BE797163, BE304820, BF239133, BE792803, AV655410, BE869379, BE250059, AA310938, BE279363, AA490809, BE293423, BE293826, BE612529, BF509241, AW139250, AI970384, AI961068, AA923298, AW027490, AI624751, T67610, BF209934, AI917354, BE881635, BE869865, AW250249, R88163, BE908253, T72363, BF514254, AA151130, BF516041, BF971860, N36317, BE907720, AA302588, AI741422, AW249482, AA573909, AV688399, AW272801, AA085764, AI052311, N24896, AW514150, AI700257, W45013, R87485, BE293649, AK025099, AF239957, and AF113795.
HWLHB64	130	1099467	1 - 1900	15 - 1914	AI768894, BE004534, AW445027, AI692721, AW451656, AA778796, BE218913, AA086137, AA827367, AI672409, BE218606, BF512195, AI240965, AA206451, BF515912, BF803236, AA643565, AI246897, AI920888, AL039827, AI669161, AI420955, AW886362, AI620406, AA011351, T81961, BF757056, AW747924, W01048, AA526319, AA342212, AW964995, BF813852, BF757054, T98421, BF754707, BE179871, BF754693, AF263462, and AB037740.
HWLED94	131	1099509	1 - 2141	15 - 2155	AU132908, BE616563, BG179837, BE563549, BE869873, BE906394, BE735172, BF205509, BE727827, BG027638, BG168740, BE546953, BG250345, BE734742, BE296124, BE889102, BE278334, BF688255, BE298950, AL535403, BG032190, BF036905, AL449873, AL449874, N47380, R51419, AA233715, BF689043, AA281397, BE386425, AA853680, N47383, AA559954, R48018, AW845348, AW963306, BF807021, BG252960, BG003101, BF899349, AW817900, AA360056, BE147292, AI817280, AI671902, U46295, H29166, AA988104, N56996, AU154347, BE858553, AA479464, BE171341, AI192374, AA659888, BF438286, AI927558, AA548397, BF510086, AI161167, AI418829, BE327906, AF300795, AK001668, and AF158748.
HKAJI55	132	1100188	1 - 1032	15 - 1046	AI751570, BE856602, BF984333, AC005821, AF134838, AB014609, AF107292, U56734, AC005821, and AC005821.
HCEPF75	133	1100345	1 - 1457	15 - 1471	BE388239, BF345885, BF346212, AW964936, BG222601, AI963082, BF346206, BF448000,

W67470, AI491829, AL036457, AA693916, AW393728, AA082338, AI282280, AI640197, F08289, AI129702, AW382744, W04784, R02486, AA043484, N39494, AW393723, AA297541, N75482, W42930, AI350453, N47632, W32485, R27136, AI274641, R40950, BE818086, W05292, AA903449, BE592075, Z41300, AA329606, BE348686, AV681500, AV681505, AV681504, AV645385, AV645390, AV645392, AV645393, AV681492, AV681509, AV681529, AV681526, AV681512, AV645336, AV645317, AV681507, AV681465, AV681525, AV681495, AV681502, AI383010, AV681486, AV681483, AV681464, AV681462, AV681487, AV681506, AV681523, AV681519, AA034469, AV681497, AV681471, R13439, AV749566, AV645358, AV645401, AV645332, AV645383, AV645377, AV645379, AV645389, AV645369, AV645380, AV645321, AV681527, AV681501, AV645400, AV645404, AV645353, AV681458, AV645334, AV681510, AV681528, AV681474, AV681514, AV645339, AV681477, AV681491, AV681472, AV681475, AW971745, AV681488, R26770, AA773671, AW779291, BE179292, AW393730, R02379, R26780, AV681513, N53823, AA595506, AA365217, R26900, R25976, N52700, R25966, AV681461, BG166813, AV645350, BF989312, BE614237, BE613349, BE612945, BE622828, BE613070, BE622931, AW885484, BE613136, BE614907, N75332, BE938236, BE614230, BE621911, H24486, BE620742, BE618811, BE614451, BE622868, BE622925, AV681531, AW883620, BE620756, AV681468, D79864, AL036403, BE764813, BF038480, BG003969, AW877209, AX015418, AR085071, AL137638, U69263, U69262, AR080280, AR091518, AR093384, AR035224, I85513, AR009152, AR009151, AR027099, AX009487, AR093392, I25041, AR093383, AR069374, I25027, AR069375, I44515, I26928, I26930, I26927, I44516, A94046, A94054, I05393, A10617, AR028792, A01323, AR034783, AX030966, I63120, AR067733, AX009486, AX029455, AR054109, AR064322, AR064323, AR064320, AR064321, A94048, A94061, AX027785, AR038307, A01324, A32110, A49045, AR038321, A83642, A83643, A70359, AX003194, AX035462, AR083151, I05430, AR019094, A92666, A92668, I49890, A92667, A92665, AR028791, AR028793, AX032758, A91752, AR018924, AR018923, A48774, A48775, AR000006, AR015960, AR015961, AR051652, AR000007, AR091393, AX008865, AX008867, AX008868, AR069413, A92081, AR069417, A92080, A92077, A92078, A92079, A91751, AX030369, AX030368, AX036660, AX036661, AR068508, AR068510, AR068509, AR091571, AR085090, AR003585, I58322, I58323, A91754, AX026821, A63067, A51047, A63064, A63072, AR031375, AR068507, AR068506, AX001323, AX001324, A85308, AR062871, AX044426, AX044425, A60213, AX033488, AX046332, AX033489, AX033490, A44171, AX027908, AX027904, AX033474, AX033486, AX033487, AX027906, AR068550, A23373, AR068551, A49700, A60207, A60208, I58669, A58521, A29109, A32111, AX004550, AR031374, I07209, I07249, A63954, AR019098, AR020199, AR020200, AR001287, AR020198, AR020197, AR029418,					
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HDPK184	123	1096306	1 - 930	15 - 944	BE264355, BF316651, BF855767, BF026003, AI949213, BF027434, BE794314, BF793601, AW440269, BE407465, AR084814, AF030335, AJ300588, and AX001017.
HDPUN70	124	1096317	1 - 530	15 - 544	AW963155, AA361063, AA535015, AA361359, AV721179, BE568285, BE766706, AR035946, AY009090, AC006510, and AC006510.
HHEPF57	125	1096422	1 - 3583	15 - 3597	AL520380, AW964307, AI861833, BF062378, AU137320, AI161343, AW967203, AW954939, AI221950, AL041826, AI271350, AI205879, AI080159, AI183981, AA868067, BG111970, AI808607, AA452138, AA931374, AA213383, AA449162, AW961428, AA927464, AU157129, AA452364, AI668878, AI669908, AI147689, BG032378, AI360990, AI360971, W84487, AA025228, AW015764, AA476491, AI743040, H51515, T08347, AI970366, W20339, Z39489, BE972558, F02802, D80443, W84564, AI634710, N92722, T08346, AA207021, AA213601, F06532, AA025991, R42398, AA328933, AA628165, F06533, AA205234, F08486, AI352260, R14674, AL041827, R37459, C02421, F04701, N92721, AI190488, BE702258, AC004142, AL442092, D49802, AF291437, and AK001991.
HPKFH60	126	1096718	1 - 2146	15 - 2160	BG166813, BF569563, BF038480, BE868728, BF125716, AL036457, BF125688, AV734107, BF347123, BF350124, BG002711, BE868584, BF373294, BF530895, BF832657, BF371464, BE839468, BE773077, BE839432, BE839471, BE839441, BE839442, BF373232, BF371463, BF373239, BE839466, BE839452, BF373231, BE839433, BF371467, BF371480, BE839434, BE839449, BF371470, BE301851, AA298644, BF371462, AI769294, BF125601, BF371499, BF757123, AV749566, BF371508, BF371495, N86321, BF989312, BF886056, U69263, AX015418, AR085071, AL137638, U69262, and A27272.
HHBFK81	127	1096722	1 - 2455	15 - 2469	AI675601, BF448710, AI927569, AI021956, AW006374, AW471286, AI826018, AV707927, BF671821, AI890341, BF435308, AI810654, AW169258, BF570055, BF431979, AW874655, AI651209, AV702209, AW953957, AW019935, BF476689, BF697787, AI972357, AI199843, BF569563, AA065199, BE301851, AI088890, AA709124, AI148316, AI653895, AI128287, N51991, AI370980, AI131110, AI128288, AA121420, AA595022, AI050847, AI097508, AI139419, BE763321, AI089965, AA167466, BE326981, BE074862, AA973858, AI131359, AW474345, AW023165, AI073942, AI131119, W67469, AA121267, AI078524, AW513406, BF589348, AW023176, AA595425, AA773728, BE220409, BG002711, AI804337, AA931243, N26805, AW021990, AW167929, AW779991, AW149152, AA514757, AW204894, AA071473, AA016282, BF894379, AW768838, W42865, AI215875, W02151, BF439864, AW593630, AI769294, AA448616, BE868584, AA953246, AI041690, AA065130, AA716632, AA569580, AI970871, AA027942, AA782744, AA971152, N57486, AA621469, AI240615, AI332533, AA693418, AI092127, AI869057, AI755001, AI217065, BE769303,

HEOAD12	117	1095888	1 - 1124	15 - 1138	N84048, N55698, N88496, AA247827, AA095641, AC008812, Z60599, Z60600, AF102850, AX026543, AR074144, AR066487, AR074136, AF039698, AJ401268, AJ277275, AJ277276, and X98248.
HEMFC27	118	1095935	1 - 2355	15 - 2369	BF307050, BE294659, BE295378, BE270183, BF306043, BF304968, BE299101, AA380349, W19499, BE832758, AW882273, AW263540, AA157229, AA225180, AW409843, AA151564, AA909334, AY009951, and AF110865.
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HFOVM21	119	1096023	1 - 5413	15 - 5427	BF965934, BF358074, AW753022, BG238877, AI621089, AW851445, AI751442, W45174, AW069755, AW339880, AI751441, AW068275, AI752628, AW604115, BF509527, BF056654, H16112, AI825217, W45078, BF749563, BF770980, BF770976, H15811, U53091, AA852615, AA852614, AI738528, BF770978, BF928722, BF770979, BF365293, AW888738, AA665807, BF328395, BG013602, BF770969, AI750918, AI752385, AI752488, BF770975, BF770962, AA316377, BF770974, BF769640, BF758396, BF758424, AI752355, BF830392, AI750919, BF753051, BF753042, AV699874, AV700783, AW166270, AF109681, AF137378, AL359064, Z50167, and Z50157.
HCE3H71	120	1096278	1 - 2030	15 - 2044	BF530859, BE047009, AI638508, AW966265, AW139057, AW149807, AI654790, BE504594, BF432418, AI419465, BF594084, BE796905, BE207202, AI569283, H51217, H29244, R85125, R90888, H29245, AA488106, AI480182, H05353, AI500178, F11386, AI873131, AI499775, AI458144, F05862, R44423, H05303, H51247, R14779, F09051, AA338672, R51725, R51648, R42549, F07470, R35934, Z40525, R19635, W22746, R49451, R85080, F02105, BF530977, BF366816, BG055804, D29763, D64009, and D64010.
HOABR66	121	1096304	1 - 427	15 - 441	AI751571, BE543881, BF800758, BF800750, BF800624, BF992740, BF800754, T11884, BF800639, BF800628, BF800636, BF800633, AF134838, AF107292, AB014609, AC005821, and U56734.
HOHBI59	122	1096305	1 - 1632	15 - 1646	BE378623, BF312766, BF340130, BE257144, AI940172, BF568328, BE695783, BF339085,

					AW949642, AW973447, D59275, D80022, AW949629, AW949656, AW959628, AV700889, AV720812, AW949643, AV723097, AV719188, AW949632, AW949633, AW949631, AW973307, D80378, AW959582, T03269, F13647, AW975621, AW949633, AW949658, D80045, D59859, AW949646, D80166, D50995, AV742001, AV742667, AV701125, AV701335, C14014, AV701166, AV701043, AV701332, AV701017, AV701248, AV701431, AV700229, AW960553, AW966534, C15076, AV720028, AV645389, AV645344, D80269, D80134, AV718440, AW966054, AW949618, AV701419, AV701154, AW975618, AV719628, AW978661, AW965177, AV681510, D50979, AV681491, AV699746, D57483, D59502, AV745080, AV701130, AV719000, AV701149, AW966041, AW959202, AV701422, AV645343, AW966013, AV720654, C14429, AW960414, AW975605, D80024, AW959597, AV701443, D51250, AV699479, AW964737, D80268, AW966050, D81026, AW949654, AV699669, AV701415, AV701344, D80164, AW973470, AW752082, AV720220, AV681529, D58253, AW949655, D80949, AW966043, D59467, AV721784, AV719913, AV701428, AW964488, AV699682, D59787, AV720791, AV681468, AV718633, A62298, AX033851, AX027925, A62300, AX047063, X67155, AX020191, A67220, AX020190, AX021518, A25909, AR070327, A84916, AX047062, D26022, AJ132110, D34614, AX047064, AR025207, A78862, AR092424, D89785, AJ302649, Y17188, AJ294956, AR018138, AX035434, D88547, X68127, AR088705, A85396, AR074141, AR066482, AB012117, AX042372, and A85477.
HNECM88	114	1095640	1 - 968	15 - 982	AW403840, BG014445, BF359557, AC018755, U71383, and AF170484.
HCQB177	115	1095827	1 - 1128	15 - 1142	BE250620, AW955280, BE740673, AI268313, AI872092, AI935261, BE736986, AI522355, AI636626, AI673056, AA242749, AW051818, AI419938, BE906649, AI990636, AW592642, AW236974, AI829149, AW591991, AA614146, AW195603, AI066551, AI005246, AI148387, BF528484, AW027831, AI189569, BF128703, AW303516, AI269298, AI635507, AI272674, AI148390, AI370747, AA983894, AA827816, AI954835, AW406541, H99879, BE464778, AA195267, BE714240, AA922166, AI242595, BE250372, AI474318, BE714202, AA582660, AA195084, AA243887, AW605414, AW844454, AA306943, N49968, AI368413, AA157973, AW373048, BE019348, BG165521, AW629707, AA643893, BE672762, AI318617, AW629706, F08850, AI307544, AW381506, AW732270, N28564, AA186619, AW772559, AI264807, AA582370, N83852, AW190338, F11182, BF509192, BF437281, BE926434, AI674609, R95152, BE831272, BE786035, AA905701, AA093791, AI915375, AW118291, AW371665, H52011, AW371696, AW452496, BG152868, AI699638, AW883177, T23908, AA310217, BF910924, BF313744, BF304983, BF309869, N52745, BF309193, BF307702, BF664275, AW581094, BF203225, BE796766, BE261201, BE382658, BE741211, BE744257, BE250199, BE297368, BE250599, BE260377, and BE261301.
HBXFI23	116	1095877	1 - 1055	15 - 1069	BG010310, AA774984, AV690857, R96439, AA093224, N87989, N88518, AA096046,

HLJBH31	111	1095303	1 - 2178	15 - 2192	AX006821, AR067731, AR067732, A58522, A91750, AX035462, AR008408, A70867, M60510, AR093385, AR016691, AR016690, U46128, I79511, AR000511, D13509, A64136, A68321, AR060133, AR087528, Z82022, U87247, AB023656, U79457, AF123263, AR060382, AR032065, X93535, and AR008382.
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HMCHH31	113	1095550	1 - 2263	15 - 2277	BE271199, AW575245, BF794609, BF797900, BE559773, BE384088, BE513826, BE270971, BF572042, BE560978, BF690655, BE275832, BE674800, BF303959, AW205367, AW402801, BF203242, AW402242, AW402928, BF305905, AW403946, BE892536, BE466652, N24246, AW968460, BF572179, N28316, AI654541, N29315, N38941, AW383418, BE729612, AA215300, AW383426, AI276242, N33010, AW383396, N20230, BF692515, AI439520, N29316, AA459158, N25452, W03476, AW383428, AA458944, AW402824, N30453, N28949, N21241, AI760983, N20533, BF896859, N72999, N20563, BG025865, AW403434, AI434284, N67502, N73074, AA837208, AI470743, AW407871, AW404443, H84381, N26470, W02963, AI864746, W02298, N46511, H84382, H98912, N35519, H99497, BF890914, AA761778, N71796, AI222330, AW978634, D80188, C75259, AV718489, D81030, D58283, D80253, AV722801, AV699447, D51423, D59619, D80210, D51799, D80240, AV719822, AV718692, AV719324, AW966531, AV720211, AV720731, D80366, D80195, AV723927, D80227, D80219, D80391, AV699550, AV720203, D80043, AV719783, AW959570, D59889, AV718800, AV718844, AV720464, AV718770, AV724520, D80196, AV699927, AW949641, AW966053, D80212, D59927, AW966062, D80038, D80193, AV719468, AW965158, AW949645, AV719557, AW949657, AV721386,

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HSXDF41	108	1094953	1 - 2297	15 - 2311	

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HWHQR2 5	102	1094777	1 - 574	15 - 588	AB043894, AF279305, and AB036931.
HNTAV78	103	1094783	1 - 717	15 - 731	AL137118, AB038269, AB041644, and AF254664.
HNTCU89	104	1094785	1 - 604	15 - 618	BF694934, AW754392, AL118512, and AL133553.
HTSHF07	105	1094859	1 - 1941	15 - 1955	BE312916, AW976020, BG027164, AA595199, BF996520, BF341222, AW516908, AW975629, AW572281, AI339553, AI041943, AI284951, AA430329, AI652283, AW594568, BG150283, AI290327, AI087248, AI961855, AW361580, AI952525, H19258, AA730621, AW839067, BF922760, AW750496, AI081192, AW166405, AA747653, BF062691, H12557, H27260, BF063323, AW751616, R55548, AI081393, R55547, H28600, AI695272, H26634, H00825, H12800, T55811, H26693, AI968594, AI367028, AA853085, BF736378, BF358075, AA430355, H42368, AI468799, AI864807, R23780, T55383, T50030, BF848831, AI783755, F22336, AW840896, AW204995, AV725294, AW963397, AV702726, AV701983, AV705545, AW965515, AV661866, AV705217, N71226, AV703593, AV707444, AV701863, AI557264, AV702189, AW954129, AW965551, AV706545, AV726916, AV709895, AV702651, AV696302, AV703159, N71729, AW951516, AW967178, AV726708, AV702148, AV728144, AI557254, AW958016, AV703710, AV726228, AI557278, AV725247, AV708948, AI541393, AF279142, AX011625, and A84916.
HISAF60	106	1094890	1 - 4377	15 - 4391	AL524621, AL521404, AL524622, AL521403, AW978618, AL529722, BG256352, BE909801, BE783472, AW953057, W60630, W29012, AA149513, BF968833, BE783711, AA149644, AW965598, AW977554, BE834598, AA306190, AA404374, AA928795, AW026671, AA733045, AI131505, AW051295, AI139050, AI144018, AW162934, BE855656, AI089282, W76344, AL571763, AI351676, W60631, AV718844, AW005213, AA146937, BF366791, AV727103, AI146486, AV724987, AV656240, AV704592,

						<p>AR060234, AR066494, AJ279014, AR023813, A81671, AR064707, AR069079, AR096545, AJ251859, AX030435, AR054110, AB026436, AR043113, AR096546, and AR064706.</p> <p>AW504558, AW403530, AW504347, C14389, C15076, AV720150, AW949630, D59467, D80164, AW975618, D81026, AW949646, AW949641, D59787, AW964532, AW973488, AW966050, AV719822, AW966054, AW966053, D80195, AV699866, AW966534, AW975621, AV718530, AW966013, AV719783, AW960465, AV720464, AV718681, D59502, AW964477, AV718800, AV701004, AV724520, AW959469, AW978661, AW949654, AV720791, AW978648, AW959570, AW965185, AW965197, AV699927, AV718692, D81030, AW966041, D80166, AW973541, D80022, AV719324, D59275, AV718931, AV718770, AV719188, D80227, AW966531, AW966062, AW978634, AW959597, AW949658, AW753053, AW965184, D58283, AW966022, D59859, AW958992, AW959799, D80391, AW965196, D80193, D59619, AW949634, D80210, AV719468, AW960553, D80240, D59610, D51423, AV699550, AW964488, D51799, D80253, AV720211, AV718844, AV718489, AV720203, AW973307, D80043, AW966065, AV718440, AV723927, AV720028, AV718633, AW959628, AW965177, AW975605, AW949656, AW966029, AW973485, AW949642, D80038, AV719557, AV720731, D80269, AV699447, AW966075, D80196, AV722801, AW973334, D80212, AW959202, AV723097, AW959062, AV699497, AW965175, D50979, D80219, AV721386, D80188, AW956434, AW965163, D59927, AW965158, C14331, D57483, AW949632, AW973474, AW964756, AV718938, AW949645, D80366, AW958993, AW975613, AW959136, AW966059, AW966030, D59889, D80241, AW960473, D50995, AW962082, AV718707, AV720878, AW973482, AV720812, AW949653, AW949631, AW949643, AW949618, D80378, AW949655, D80045, AW949629, D80251, D51022, AW949633, AW949657, AW959582, AV741187, AV741198, AV741191, AW966043, D80024, AW177440, AA305409, AW962245, AV752082, AW973447, AV700159, AW960454, AW973330, AW956397, AV700229, C14429, AW964737, AV738340, D52291, AV700889, AW966032, AW378532, AV699652, AA305578, D51060, AW966023, C75259, AV720533, AW960564, AW960504, AV720654, T03269, AV744662, AW960532, D80522, AW178893, AV699669, AV701125, AW975623, AW962395, AV701166, AV742430, AW960570, C14014, AV699746, AV742001, AV742667, AW964468, AW179328, AV719758, AV701335, AV701043, AV701332, AV701017, AV701248, AV742048, AV701431, AV701149, AW965176, T11417, AV719000, AV701154, AV701422, AV681510, AV681491, AA514188, AW973490, D80248, AV719049, AV744690, AV699682, AV742022, AW753067, AW973465, AV720220, AV701419, AW753064, AV719628, D51250, AW369651, AV645389, AV645344, AW352158, D80134, AA514186, AV701443, AV744004, D58253, AL356276, AR018138, A84916, AR070327, A62300, AX027925, A1132110, A62298, AX033851, AX047063,</p>
HNGEO79	101	1094775	1 - 783	15 - 797		

HDAAN31	100	1094668	1 - 862	15 - 876	AA620978, AI652117, Z99396, AW979232, AW973270, AW969885, AW979204, AW973202, AW973213, AW976515, AW975976, AW979165, AW976510, AW975975, AW969884, AW970113, AW972943, AW973770, AW970860, AW969759, AW979083, BF592735, AW971732, AW976982, AW970587, AW975876, AW973650, AW979175, AW979064, AW969637, AW975938, AW975138, AW979002, AW975966, AW975968, AW972762, AW979116, AW972884, AW973987, AW975910, AW975943, AW973805, AW973164, AW971964, BF588769, AW974379, AW969752, AW973967, AW975933, AW973207, AW979176, AW973104, AW969782, AW970868, AW975959, AW973209, AW975971, AW973219, AW972440, AW979081, AW973167, AW975162, AW973946, AW975261, AW971367, AW979294, AW975229, AW979085, AW979228, AW979220, AW973190, AW974915, AW973088, AW975173, AW975955, AW974107, AW969861, AW979219, AW969839, AW972417, AW973271, AW975928, AW975964, AW971387, AW979090, AW969753, AW973997, AW975106, AW975168, AW975025, AW975031, AW975914, AW975922, AW975172, AW975387, AW969913, AW973200, AW975691, AW979078, AW975157, AW975174, AW969815, AW975022, AW973829, AW969719, AW975986, AW971096, AW973150, AW979095, AL038837, AW969661, AW973168, AW973761, AW975944, AW969797, AW975276, AW973738, AW975637, AW975917, AW969673, AW969751, AW975594, AW975953, AW976045, AW970877, AW975277, AW974784, AW975935, AW970969, AW979106, AW975181, AW971975, AW969755, AW973762, AW975950, AW969646, AW971938, AW973793, AW975930, AW973183, AW973986, AW973815, AW975029, AW979088, AW973778, AW975931, AW969794, AW970945, AW975046, AW975956, AW975105, AW979068, AW973241, AW969804, AW979144, AW971937, AW970921, AW979194, AW971408, AW975978, AW976511, AW979174, AW970958, AW973171, AW969766, AW969895, AW971963, AW973101, AW973812, AW975981, AW975058, AW975947, AW979221, AW975033, AW973132, AW970947, AW972806, AW975923, AW975429, AW969662, AW971950, AW971965, AW979076, AW970588, AW975632, AW973788, AW974981, AW976042, AW979127, AW969682, AW975988, AW971984, AW973103, AW973789, AW973785, AW974139, AW975032, AW970097, AW973824, AW971974, AW974787, AW979284, AW975904, AW971731, AW974806, AW975130, AW973810, AW969685, AW972156, AW974823, AW974658, AW975965, AW970997, AW973165, AW969668, AW970062, AW973808, AW972419, AW973116, AW973806, AW973823, AW970044, AW971001, AW975154, AW973771, AW975915, AW971985, AW972296, AW969783, AW969803, AW972784, AW974790, AW970933, AW973772, AW975042, AW973985, AW979197, AW969714, AW975987, AW971129, AW971403, AW975015, AW976044, AW979142, AW974785, AW975649, AW975002, AW975177, AW973784, AW979178, AC007023, AX046357,
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HLKAB61	96	1094201	1 - 503	15 - 517	AW024960, AI479960, AI052585, AA830014, AW205456, AA398256, AI859053, AW137104, AI873741, AI796467, AA225625, AI159823, AA811798, AI718153, N25872, AI969940, AI382438, AI311717, AW088488, AI187156, AI278434, AI088449, BF590892, W92243, AA761003, AI631818, AA146624, BF062897, AI494095, AI452434, AI066392, AI245054, AA535733, AI190865, R59937, AA158516, N30762, BF002429, AA634216, BG168579, AA564062, AI139986, AA135576, BE615149, BE615178, W70120, BF111558, W99328, AA399431, BE615246, BE000703, BE001017, BE614950, BE616347, AV711597, AA226160, N45043, BE616626, BF346442, AA143555, AI906964, BF084997, AA190958, AA034276, AI708811, BE925851, N40580, BE814127, W69970, H94758, AW338143, BE814776, AL049715, AF315356, AF308601, and M93661.
HFVZB56	97	1094512	1 - 1510	15 - 1524	BF994524, BF994526, BF994521, BF998660, BF369110, BG006771, AA298425, BF998991, BF368046, and AF206329.
HAGFI46	98	1094598	1 - 1317	15 - 1331	AV709803, AW954538, H23116, R19556, R71701, BE009844, T27055, AA223802, BF850808, BE150550, AA349536, AC008039, and AF196976.
HYAAM4 3	99	1094663	1 - 976	15 - 990	BE311916, BG164873, BG231605, AW511388, AI990505, AW295528, BF061905, AI761169, AI968491, AW467179, AW448991, AW264297, AI825483, BF899230, AW467175, AW467180, BF896246, BE812455, AW236598, AV756474, BE924973, and AB024718.

HELGM30	95	1093937	1 - 6568	15 - 6582	<p>AW956434, AW964488, AW949641, AW962082, AW949656, AW949654, AV699927, AW949642, AW959202, AW973470, D81030, AW973468, D81026, D80045, AW179024, AW176467, AV739950, D80212, D80196, D80188, AV718681, D80219, AW978633, AV742732, AV701154, AA305578, C15076, D80038, D80269, AW352171, D59610, D80366, D57483, AW377676, AA305409, AV744004, D51022, AV744006, AW966043, D50979, AW352170, D50995, AW177731, D59889, AW178907, AW951169, AW753041, AV721784, AW178762, AW962245, AW179019, AW973447, AV742022, AW966023, D80024, AF039686, AF118670, AR034800, AF081916, A84916, AX047063, Y17188, A62298, AR018138, AX047064, A62300, AX047062, A82595, AX033851, AJ302649, Y17187, AR070327, X82626, D89785, A30438, AF058696, A67220, A78862, AX028130, X67155, AX020191, AX027925, AB028859, D34614, D26022, AJ132110, AX021518, AR008278, Y12724, AX035434, AX020190, A25909, D88547, AR087649, AR025207, A94995, AF260572, AR008443, I50126, I50132, I50128, I50133, AR066488, U79457, AR016514, AR060138, A45456, AR066487, AR074136, A26615, AR052274, D50010, AR074139, Y09669, AR074545, AB012117, AR060385, AB002449, I18367, AX015396, A43192, A43190, AR038669, AJ294956, A85396, AR074141, D88507, AR066482, AR088705, AX042372, A44171, AJ287395, A85477, I19525, A86792, U46128, D13509, AR016691, AR016690, AR093385, X93549, AR066490, AR008408, AR060133, A63261, A70867, AR062872, AR032065, AF123263, AR091537, A64136, A68321, I14842, AR054175, and AR008382.</p> <p>AL534116, AL521556, BE799405, AU117881, BE796852, AU125856, BE618804, BE618292, BE799308, AW167760, BE799155, BE871095, BE794749, BE869883, BF219975, BG259354, AW962203, AW886515, AW365110, BE251414, AW955201, AW953036, BE618274, BE170498, BE296419, BE563317, AW960058, BG258902, BE296266, BE294341, AI304882, AL039240, BE293232, AW960968, BE263370, BE253347, AA173227, AW955661, BE618794, AL048118, AI161133, BF203822, BE253486, BF222454, BE778159, AA918989, BF037372, AW842412, BE269763, AI831411, AI916267, AI142864, AA227614, AW842323, AI990905, AW249103, AW248406, AW270995, AI935433, AA640948, AA548712, AI160268, BE673909, AW248670, AI582201, AW778831, AW842385, AW842333, AI127168, AI768288, AI692762, AA788923, BF435484, AA227913, BG056383, AI559620, AI521614, AA421940, AI051695, AI590586, AI097334, BE326236, AA411274, AI271548, AI343458, AI952851, AI681495, AI051716, AW592477, AW795474, AI479202, BE265320, AI131409, AW473159, BE251845, AI950822, AI055832, AW300713, AA700761, AA707235, AI380833, BF813122, AA971227, AW316742, AA063635, AW338425, BF437556, W46855, BE258943, BF220047, T08335, AI146822, R54648, AW383943, W95244, AW157703, AA836747, BE156124, AA693411, AW300605, BG055076, AW842358, AW205519.</p>
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HATCZ07	92	1093241	1 - 1139	15 - 1153	AL524519, BF792657, BE561100, AL524518, BE348474, AA446779, AA493561, BF851237, BF933078, AA115893, AA446757, AV681737, AW378165, R06836, AA216146, AA197081, AA115848, BF247394, BF516391, R18502, T46848, T39504, H28550, AA297984, BF922563, BF928739, AV696103, N99420, AA443398, AA834902, AA197090, AA621288, AA659631, W67535, AA723129, AA884255, AF132811, AF061260, AB021966, AB021964, AB021965, AB021967, AB017560, AB017558, AB017556, AB017555, AB017557, AB017554, AB017559, and AB017561.
HDPXT07	93	1093488	1 - 2568	15 - 2582	BG258479, BF796825, BF821423, AA534416, BF821426, BF879214, BE463469, AW886669, BF368287, BF794125, T91577, T91625, AW299992, AI689842, BF590018, BE818151, and AA782975.
HMCHG54	94	1093752	1 - 1951	15 - 1965	AV733801, BF196710, AI093369, AW292321, AA972431, H97311, AA746376, N40174, AA130392, AA286750, AA287684, R71586, AW889046, BF366889, R71568, R71587, H03136, AW889047, H03946, R71567, AI471079, BF366460, BF366567, BF366574, BE766489, N54926, AA365025, AA451915, AW964468, AW975618, AW9666389, AW964532, AW949645, AW966330, AV742048, C14331, AV724520, AV741220, C14429, AV738340, AV718489, AW360811, AW752082, AW753053, AW177440, AW375405, AW375406, AV699550, AW177501, AV720791, AW177511, AW178893, D59467, AV741221, AV718692, T03269, AW179328, AW352117, AW973423, AV741191, AW366296, AW973541, AV742430, AV726921, AW360844, C14389, AV701004, AV727978, AW973490, AW377671, AV758213, AW753067, AW360817, AW966053, AV701125, AV702035, AV701166, AV701149, AW966029, AW966075, AW966065, AW960465, AW973334, AW966531, AW978634, AW378534, AW966534, AW179332, AW966560, D58283, AW966022, AW377672, AW179023, AW178905, AW959799, AW966059, AW966013, D59859, AW964766, D80022, AW966041, D80166, AW973474, D80195, AW975621, AW978648, D80193, D59927, AW975613, D51423, D59619, AW978661, D80210, D51799, AW965163, D80391, D80164, D59275, AV720533, AW960553, D80240, D80253, AW966030, AV719822, AV744690, AW966054, AV720203, AW964756, AW966050, AV719188, AW973307, D80043, D59787, BF349328, D80227, AW966062, AV719324, AV718440, AV718938, AV719783, AV720028, D59502, AW959597, AV718633, AW959628, AW960473, AV742001, AW965177, AV742667, AW975605, AW959570, AV719468, AV718800, AW965185, AW965197, AW965196, AW973485, AV718707, AW965184, AW973488, AW965175, AV720211, AV718931, AV701335, AW962395, AV720878, AV718844, AV720464, AV719557, AV718770, AV720731, AW973482, AV699447, AW378532, AV744662, AV701043, AV701332, AV741187, AV701017, AV701248, AV741198, AW958992, AW958993, AV701431, AV722801, AV723927, AV699866, AW959136, AW956397, AW959062, AW964477,

					<p>AW378532, AW949645, AV701004, T48593, AW375405, C15076, AV699866, AA305578, AW966330, D80269, D57483, D59610, D80038, D80366, AW973447, AW178775, AV70220, AW177501, AW966043, D51022, AW177511, AW377671, D50979, AW949653, AW949631, AW949643, D59889, AW949618, D50995, AW949655, AW962245, D80024, AW956397, AW966023, AW959469, AW964737, AW966032, AW959582, AW960532, AW965158, AW949629, AW949646, AW949633, AW949632, AW949658, AW949657, AV742022, AV701443, D81026, AW753041, AV701123, AV701130, AV701419, AW973445, AV744662, AV701154, AV741187, AV741198, AV741191, AV701422, D80378, D51060, AW973490, Z59035, AR070327, A62298, A30438, D26022, A84916, AX047063, U46128, A67220, Y17188, AX047064, AX033851, AJ302649, A78862, A62300, AX035434, A25909, X67155, D89785, AR018138, X82626, AX027925, AJ132110, AX047062, AF058696, D34614, AX028130, AX020191, AX021518, AX020190, AR008278, I50133, Y12724, A94995, AB028859, D88547, AR087649, AR074139, AR025207, D50010, A43192, A82595, A43190, AR038669, AR008443, AR060138, AJ294956, I50128, I18367, I50126, I50132, AR066488, AF260572, AR016514, Y17187, X68127, A45456, Y09669, A26615, AR052274, AR060385, AR074545, AB012117, AR062872, AB002449, AR066487, AR074136, AR066490, AX015396, AR008408, A64136, A68321, AJ287395, A63261, A85396, AR074141, D88507, AR066482, A44171, AR088705, AR016691, AR016690, A85477, AR093385, I19525, A86792, A70867, X93549, AX042372, D13509, AR060133, AR091537, I14842, AR054175, U79457, AR032065, AR008277, AR008281, AR008382, AR050680, and AF123263.</p>
HNTDM29	91	1092545	1 - 651	15 - 665	<p>H91914, BE293969, BG251102, AW409699, BF305486, AW005786, BE781884, BE255612, BF127903, AI809532, BF799167, AI453056, AA778576, AA683020, H91861, BE041585, AI359703, AI042586, AW205217, AW971745, AW392670, AW804686, AW604723, AW861889, AW858455, BE695785, AW363220, AW858526, AW861944, AW858525, AW384394, AW604726, AL119396, AW877209, AL119319, U46347, BE705903, AL119355, BE705906, U46349, AL119457, AW577135, BF868687, AL119324, AW372827, AL119483, Z99396, U46351, AL119497, AL119443, AL119484, AL119363, AL119391, U46350, U46341, AL119444, AL119341, BE705905, BF868697, AL119439, BF868684, AL119522, U46346, AL119335, AL119401, AL134528, AL134538, AL037205, BE705904, AL119399, AI142137, AL134527, AL119496, AL119418, AW861954, U46345, AL042614, AL042984, AL043029, AL043033, AL042542, AL042544, AL043011, AL134542, AL042450, AL043019, AL042965, AL042975, AL043003, AL119464, AL042551, AF307079, AB020628, AF111097, AF111098, U72487, AF081144, AF081146, U78105, AF081145, AF081147, AF111099, AB026436, AR080280, AX046357, AJ251859, AX030435, AJ279014, AR060234, AR054110, A81671, AR066494, and AR069079.</p>

					<p>AI093294, AI949100, AW770899, AI590664, AA775800, AI192462, AI127266, AA242879, AA642345, AA191263, AI341959, AA113413, AA993392, AI144376, H49173, AU153039, AI356709, AI753033, AI143178, AA861905, AI689142, AA346616, AW193900, BF090943, AU147228, AI807754, AA961065, AU148682, AI085783, BF058547, AA196274, AA190829, AI090590, AI754727, AV656934, AW069540, T98172, AI753670, T69976, AA029313, AI799528, AW954449, AA187246, AA565674, AI092999, AA376600, AA633645, AU151366, AA653390, AI081054, W19187, AW664104, AA614456, AI192579, AI865846, AA486286, AI384069, BE219810, AA555181, AI907704, R84542, AA242813, W42499, W86626, AW151647, AI302581, AA678601, AA113267, AI040375, AA252153, AA101407, AA180337, H83604, AF200348, D86983, and AR020614.</p> <p>AL531425, AI022193, AW139382, AA484435, AW470952, AL118990, AI499684, AI936673, H94666, AA828353, AI500265, BF002873, W25099, AI695809, T80683, AA773578, AA865715, T81364, AA703007, T72188, T62061, T72967, AA700860, T68210, T68646, T71737, AC010642, and AK027222.</p>
HLDAB75	89	1092493	1 - 1696	15 - 1710	
HMSFP50	90	1092497	1 - 1833	15 - 1847	<p>BE621832, AL519962, BE895345, BE965639, AI817123, AI393356, BE621509, AI148200, AA693721, AA746047, BE833124, AL519963, AI432698, AA341942, BE294962, N36002, T03269, AW975618, D51250, AV738340, AV718489, AV724520, AV742048, AV718692, AW960553, AV699479, C14331, AW964468, AW966041, AW973541, AW752082, AW753053, AW177440, AV723927, AV741221, AV744690, C14429, AV699927, AW178893, C14389, D80164, AV719822, AW360844, AW966053, AW951169, AW973488, AV699550, AW360811, AW964766, AW179328, AW966029, AW966075, AW966065, AV742732, AW960465, D51799, AW965163, AW965175, AW973334, D80166, AW966531, AW978634, AW966534, AV719468, AV718800, D59619, D80210, AW973330, D80240, D58283, AW966022, AW959799, AV719324, AW966059, AW966013, D59859, AV720211, D80022, AW973474, AW966054, D80195, D81030, AW949656, AW975621, AW978648, AW949642, AW975605, AW753067, AW975613, D59467, D51423, AW978661, D80391, D59275, D80253, AW966030, AV720791, AV720203, AW964756, AW966050, AV719188, AW973307, D80043, D59787, D80227, AW966062, AV718440, AV718938, AV719783, AV720028, D59502, AW959597, AV718633, AW959628, AW960473, AW965177, AW959570, AA305409, AW965185, AW965197, AW965196, AW973485, AV718707, AW965184, AV718931, AV720878, AV718844, AV720464, AV719557, AV718770, AV720731, AW973482, AV699447, AW958992, AW958993, AV722801, D80212, AW959136, AW959062, AW964477, AW956434, AW964488, AW949641, AW962082, AW949654, AW964532, AW959202, D80219, AV742001, AV742667, D80193, D59927, AV701125, AW960454, AV701335, AW178762, AV701166, AV720533, AV742430, AV701149, AV701043, AV701332, AV701017, AV701248, AV701431, D80196, D80188,</p>

					<p>AV719000, AV720150, D81026, AW964468, F13647, AV701130, D51250, AV720607, D80268, AV720533, AV744690, AW177440, AV719727, AW951169, AA305578, AV718674, AV745080, AW178775, AV700622, AW960474, AW960414, AV700159, AW949630, AW964532, AV719913, AV701428, AV720220, AV701415, AV701357, AV701344, AW378532, AV681529, AW973470, C14227, D58253, D80522, AV720755, AW966049, D80949, AV721784, A62300, A62298, AX033851, A84916, AJ132110, AX027925, AR070327, AX047063, AX047064, AR018138, AX021518, X67155, Y17188, D26022, AX035434, A25909, AX020191, A67220, D89785, A78862, D34614, AJ302649, AX020190, AX047062, D88547, AF058696, X82626, AR008278, AX028130, AB028859, AR025207, AR092424, AR077702, AJ294956, AR087649, Y12724, AF260572, AR074545, AB012117, AX015396, X68127, AJ287395, A82595, A85396, AR074141, AR066482, A44171, AR088705, AX042372, A94995, A85477, I19525, A86792, AR060385, AB002449, X93549, AR008443, I50126, I50132, I50128, I50133, AR066488, AR016514, AR060138, A45456, A26615, AR052274, AR074139, Y09669, A43192, A43190, AR038669, AR066487, AR074136, AR066490, A30438, AF135125, I18367, AR054175, D88507, I14842, X64588, D50010, Y17187, AX035429, AX035428, AX035426, A63261, AR008277, AR008281, AR008408, AR091537, AR062872, A70867, AR093385, AB033111, AR016691, AR016690, U46128, I79511, D13509, A64136, A68321, AR060133, AR064240, AB037923, AR087528, U87247, U79457, AF123263, AR032065, Z82022, AR060382, X93535, and AR008382.</p>
HEPNE19	88	1092413	1 - 3801	15 - 3815	<p>AU123821, BE741610, AU126215, AU130266, AU134393, AU131275, AU124913, AU117829, AU121408, BG117370, BE746370, BF033071, BE745284, BE741318, BE748350, AW084503, BF204988, AW852329, BF309081, BE294814, AW377564, AW852052, AW852060, AA461165, AA152257, AI375044, AW389528, AW377487, AA461339, AA029866, AW852068, AU129192, AW852122, AA248351, AW389511, AW377595, AW852063, D83862, AA455307, BE748820, AW852057, AW852272, AA164857, AI160848, AA155941, BF852017, AW753017, AA450170, AI092729, AF147073, AW389530, AW369027, AA242840, BF851989, W70089, AI186513, AA669825, W86533, W04807, AA085633, AA488536, BE245341, AW367947, AA179761, AA489307, AA187378, BE695683, BE700948, AW878062, D31085, AA295358, AA083299, AI952379, AA852329, AA343268, T94048, AA375799, AA310111, AA101439, AA340431, BE695701, AA340816, AA339155, AA340909, AA190681, BE695839, AA374075, AA232462, AA639852, AA329739, BE834287, H01758, AA375517, AA242739, AA375708, AA375223, AA346892, BE932840, AA328932, W42650, AA248840, AA359153, BF361156, AA716174, AA359079, AA488403, AA376570, AI989581, AA375091, AI754458, BF032434, T92956, N55866, AA196125, AA375146, BF871732, AU152255, AA489491, BF091002, AA216443, AI804151, AA495727, AU144555, AI686975, AA347642, AI129793, AW262139, AA242948, AI352683,</p>

					AX021518, I50126, I50132, I50128, I50133, X67155, Y17188, D26022, AX035434, A25909, AR092424, AX028130, AX020191, A67220, D89785, A78862, D34614, AJ302649, AX020190, Y12724, AR016514, A94995, X68127, AR060138, A45456, A26615, AR052274, AR066488, Y09669, AR074139, A43192, A43190, AR038669, I14842, AR008443, AX047062, AR066487, AR054175, AR074136, D88547, A30438, Y17187, A63261, X82626, AR008277, AR008281, AR093385, D50010, AR091537, AR062872, A70867, AR016691, AR016690, U46128, AR008408, AR016808, AR025207, A64136, A68321, I79511, X64588, D13509, AR060133, I18367, AX014811, and AF123263.
HMEBH77	87	1092363	1 - 963	15 - 977	BE144811, AW966053, AW975618, D59859, AW978634, AW960465, AW959799, AW949633, D80391, AW966534, AW966022, D59787, AW966531, AW964488, AW966041, AW959570, AW978661, AW973541, AV719822, AW966054, AV718692, D80196, D59275, AV718440, AV720028, AW965158, AV720203, D58283, AV718770, AV699447, AW966013, D80253, AW949654, D80022, AW973307, D80166, D51423, D81030, AW966043, AV719783, D80195, AW975621, AW978648, AV720464, D59619, D80210, D51799, AW960553, D80240, AW949641, AV720791, AV718489, D59927, AW966050, AV719188, D80043, D80227, AW966062, AV719324, D59502, AW959597, AV718633, AW959628, AW965177, AW975605, AV718800, AW973485, AV720211, AV718931, AV718844, AV719557, AV720731, D80024, AV722801, AV723927, AV724520, AV699550, AW949656, AV699927, AW949642, AW959582, AW959202, D80212, D57483, D59889, AW973334, D80188, D50995, D80219, D80366, AW964756, D59467, AV718938, D80269, D80038, AW956434, C14389, AW966029, AW966065, D50979, D80193, AV721386, D80164, AW949629, AW949632, AW965163, AV719468, AW949645, AV718707, AW949657, D59610, AW958992, AW958993, AW949653, AW949646, AW949658, AW949631, AW949643, AW949618, AW949655, D80378, AW959136, AW959062, AW964477, C14331, AW959469, AW962245, AV720812, AW965197, D51060, AW966075, C14429, AV723097, AW973447, AW973474, AW964737, D80241, D80045, AV700889, AW956397, C15076, AW965185, AW965184, AW962082, AW973488, AW960504, AW960564, AW960570, AW965175, AW973482, T03269, AW975613, AA305409, AV720878, AV700229, AW960473, AW965196, AW973330, AW966059, C75259, AV720654, AW752082, C14014, AW753053, AV645389, AW966023, AV701004, AW975623, D51022, AW960454, AV742001, AV742667, AV701125, AV701335, AV701166, AV742430, AV701043, AV701332, AV701017, AV701248, AV742048, AV701431, AW178893, AW965176, AV699669, AV718530, AV719628, AV701149, AV699746, AV681468, AW966030, AW960532, AV699682, AW973465, AV719049, AV701419, AV701154, AV701422, AV681510, AV681491, AV645344, AV645343, AW966032, D80134, AV701443, AW964766, AW966560, AW179328, AV699479,

H2CBE03	86	1092357	1 - 778	15 - 792	<p>AR080573, AR080581, AF113130, AF035013, AY005133, AL353572, and AL353572.</p> <p>AU130934, AU124737, BE548429, AU130717, AA307070, AU133832, BE315390, AU124892, BF686099, BF683792, BE393022, BE543330, BF734606, AW795299, AV718707, D80268, AW964468, D80269, AV719049, AW975618, FI3647, AW960514, AW962245, C14389, AV700229, AW960474, AV700086, AV700313, C14014, C06015, AW973424, AW973334, AV701123, AA305578, AV720150, D80522, AW966388, AW966075, AW966065, AW753064, AV700357, AW959570, AW959597, AV718489, AW369651, C14227, D50995, AW966368, AW177440, D51022, C14331, D80391, AV721386, D80248, AV720203, D59787, AW959202, AW178986, AW966386, AV718770, AW956434, AW966053, D81026, AW964488, AW962082, AW966029, D81111, AW960465, AW966531, D58283, D59619, AW966059, AW978634, AW966534, D80210, D80240, AW965153, D50979, AA514188, AW973474, D80195, AW973465, AW966022, AW978642, AA305409, D80196, AV720878, AV718844, AW959799, AW973482, AW966013, AW958992, D59859, AV723097, AW949656, D80022, AW973307, D80043, AW966041, AW949642, D80166, AW960473, AW965177, AW975621, AW978648, D59927, AW965175, AW975613, AV719557, D59467, D51423, AW978661, D51799, AW965163, D80164, D59275, AW960553, D80253, AW973541, AW959136, AW966030, D80038, AV719822, AV699550, AV720791, AW966054, AV718692, AW964756, AW966050, AV719188, D80227, AW966062, AV719324, AV718440, AV719783, AV718938, AV720028, D59502, AV718633, AW959628, AW975605, AV719468, AV718800, AW965185, AW965197, AW965196, AW973485, AW965184, AW973488, AV720211, AV718931, AV720464, AV720731, AV699447, AV718908, AW958993, AV722801, AV723927, AV724520, AW959062, AW964477, AW949641, AW949654, AV699927, D80212, D59653, AV692290, D81030, D80219, AW973473, D51060, D80188, AW960483, AW966385, AW949633, AA514186, AW959469, AW966398, AV720812, D59889, AV718530, Z21582, D80439, AW966399, AW753067, AW964532, AW966330, C15076, AW966331, AW966333, AW966043, AV699746, D59610, AW966369, AW973490, D57483, D80366, AW973445, AV701216, AW966332, AW966397, AW960532, AV720151, AV720533, AW949653, AW949631, AW949643, AW949618, AW949655, AW950578, D80193, AW966400, AV720616, AW966032, AW949498, AW956397, D80045, AW949629, AW966342, AW949632, AW973447, D80024, AW966378, AW964737, AW960454, AW962395, AV720729, AW973330, D80247, AW959582, AW966380, AW965158, AV699866, AV701839, AW952852, AW949645, AW949646, AW949658, AW949657, AW952839, AW966023, AV654329, AV655880, D79997, L76158, X95351, AX033851, AJ132110, A84916, A62300, A62298, AX027925, AR070327, AR018138, AR008278, AF058696, AX047063, AX047064, AB028859, AR087649, A82595, AR060385, AR077702, AB002449,</p>
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HTEEF19	84	1092204	1 - 2303	15 - 2317	NT3970, BF447084, AI755166, BF061416, AI920880, AI925857, AW072087, BE044048, BF754597, AI215589, AI354891, AA659919, AI271925, AI635236, AI248143, AA419156, R74346, AA419119, AA477132, AA326493, AA904372, AA477379, R74443, AI187966, BF941777, T33009, N91382, R72137, BF928844, Z45408, Z41119, AA666033, AA385063, BF448827, AA349368, BF196368, AW953123, BF941802, BF940219, R72086, AI932928, BF849745, BE707723, AA091149, BF090935, AI040362, AI264069, AA464975, AI346211, R55970, N99981, AA133535, AI276142, BE466035, W79505, AW206373, AA115760, R08200, AI679070, AI934793, BE936729, H44191, BE936715, AI953019, R55612, AI677914, T87783, AA465043, AC005237, AL050143, and AK025374.
HEEAE73	85	1092290	1 - 2145	15 - 2159	BF966810, AW512865, AW629068, BF036927, BF444998, AV708679, BF966262, BF432984, AW005816, AA156950, BE390534, AW964432, AV706756, AA923729, AA156858, AI690584, AI078826, AI272207, H06189, BE219776, AW772529, R43993, AI245121, AI301968, AI458554, AI479563, H17616, AA987922, H52474, AI916500, AA775361, BG007392, BE221274, H08950, BE838084, AA344450, BF889177, AW207510, H17727, BF056145, H52475, AA319839, AA318767, H06457, AA758116, N77061,

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HDQGZ89	81	1092002	1 - 1459	15 - 1473	BG109779, BF974210, BE544126, AW993633, AW977048, BF797443, AA828379, BG166479, AA837189, BG177582, BF697657, BF700161, AA721432, BF447259, AI826972, AW451560, BE695785, AI134902, AK025338, AJ251859, AX030435, and AR043113.
HWAFW3 9	82	1092095	1 - 574	15 - 588	AV689583, AW503911, AU140225, AW502888, T27535, AC020923, AC009073, AF106037, AF222340, AB011097, AF183569, AF148323, AF148324, and AB047552.
HE8UL90	83	1092181	1 - 764	15 - 778	AW973541, D80247, AW966029, AW964468, AW966030, AW959570, AW978661, D80022, AV719822, D51060, AW949646, D80195, AW966389, D58283, AV718692, D80043, AW959628, D80391, D59787, AW965196, AW965184, AV720150, AW965163, AV699550, AV718844, AW966342, AW973447, AW952852, D80196, D59467, AW973334, AW966531, AW964477, AW949632, AW966054, D80522, D57483, AW966075, AW966065, AW960553, C14389, C14331, AW973488, AW973482, AW966388, AW962082, AW964756, AV718440, AV720028, AW960473, AW965177, AW960483, AW973485, AV654329, AW958992, AW975618, AW966059, AW952839, AW973490, AV719468, AV718800, AW966053, AW964737, AW975621, AV720729, D80439, AW959062, D80253, D80302, AV719783, AW973474, AW973465, D59889, AV720464, AV719049, AW960532, AW960465, AV720151, AW966013, AW966399, AW978634, AW966534, AW966050, AW966369, D80166, AW966330, AW966022, AV720616, AW975613, D59619, AV699447, AW959799, AW966333, D80210, D80164, D80240, D59859, AV719324, AW966041, D59502, C14014, AW973473, AA305409, AW973445, AW950578, AW965185, AW965197, AW978648, AV720211, AW962395, AW966368, AW966385, AV718770, AW966332, D51423, AW966397, AW949498, AV723927, AV724520, D51799, AV699866, D59275, AW949653, AV701839, D80038, AW949656, AW949654, AV720791, AW949658, AW949631, AW949643, AV718489, AV720203, AW949618, AW949642, AV719188, AW973307, AW949655, D80227, AW966062, AV718938, AW959597, AV718633, AW975605, AW966378, D80366, AV718707, AW966386, AW965175, AW966400, AV718931, AV720878, AV719557, AV720731, AW966331, AW966398, AV655880, AW966380, D51022, AV700229, AW958993, AV722801, AW959136, AW956434, AW949500.

AA573374, BG167189, AA039871, BE819588, BE746649, AA248612, T78267, AA190743, AA297663, AW672898, BF371522, BE144810, BE839410, BE838222, AA298500, Z24831, AW873503, AA429894, AA375075, AA029405, BF330449, AA374275, AL046932, AA375387, AA070973, AA251942, Z28837, T12175, AA705655, AA905600, AI559477, AI124100, AI199683, AA558142, AI810140, AA609693, AW245076, AA286738, AA095263, AA702826, AA026161, N88589, W25088, AA070974, AI493931, AW176118, BF365700, AI620967, AA041470, AA758806, BF371516, W74455, AW391879, AA845873, AA248318, BG005404, D51799, AW966053, AW975618, AV718800, AW949629, AV720203, AW966531, AW978634, AW966534, D59859, C14429, AW966062, D80391, D59927, D58283, AV719468, AV699927, AV718770, AW966013, AW973307, AW966041, D80166, D80196, D80195, AW975621, AV718489, D51423, D59619, AV719188, D80227, AW978661, D80210, AW959628, AW960553, D80240, D80253, AV719822, AW966054, AV718692, AW966050, D80043, AV720731, AA128310, AV724520, AV719324, AV718440, AV719783, AV720028, AW959597, AW965177, AW959570, AV722801, AV720211, AV718844, AV720464, AV719557, AV699447, AW248057, AV723927, AV699550, AW949641, AW949656, D80212, AW949642, AW959202, D80188, AW949633, D80219, D80269, D81030, AW949632, D80038, AW964488, D59889, D80193, AV720791, AW949654, AW949655, AW965158, AW949653, AW949645, AW949646, AW949658, AW949631, AW949643, AW949618, AW949657, AW960465, D80366, AV718633, D57483, AV700889, AW975605, AW966043, AV720812, D59502, AV721386, AW975682, D80022, AW959062, BF110739, D50995, D59275, AV723097, AW973447, D59610, C14331, AW959582, D80045, AW973485, AV718931, AW959799, AA905643, AW959469, D80378, AW966022, D59787, D50979, T03269, AW956434, AW978648, D80241, AW964737, AW964756, AV719628, AW973541, C75259, D80024, AV720654, AV718938, D80164, AW973334, C14014, AV718707, AI271741, AV742001, AV742667, AW965176, C14389, AV701125, AV701335, AW966029, AV701166, AV701043, AV701332, AV701017, AV701248, AV742048, AV701431, AV699682, AW966065, AV700229, AA305409, AV699669, AW965163, AW958992, AW964477, AW958993, AW959136, AW962245, AV719049, AV699746, C15076, AV701419, AV742430, AV645389, D59467, AV701154, AV681510, AV681491, AW965197, AV645344, AW966075, AB023209, AK025843, AK000845, AF205078, AF077041, AF151909, AL050093, A62300, AX027925, A84916, A62298, AJ132110, AR070327, AX047063, AX033851, AX047064, X67155, AX020191, A67220, D89785, A78862, AX020190, AX021518, A25909, AX035434, D26022, Y17188, AX047062, AR018138, D34614, AJ302649, D88547, AR025207, AR092424, X82626, AF058696, AR008278, AJ294956, AB028859, AX015396, AF260572, AX028130, AR087649, AB012117, X68127, Y12724, AR088705, AR074545, AJ287395, A85396, AR074141,				
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HCSF90	78	1091570	1 - 4792	15 - 4806	AI526737, AI527357, AI520133, AI527319, AI526770, BE531120, BE615116, BG249131, BG167313, BE900315, AL039400, BE884344, BG249782, AV753572, BF308572, BE893950, BE542224, N25612, AI553834, BF663432, AI134574, AW512989, AL042597, AI871719, BF058945, AA453727, AU127908, AW499567, N36098, BF339648, BE278930, BF057402, BF061439, BE392204, AA486431, BF924312, AU145864, BE439990, BE615650, AA662205, AI140127, AI457676, AI811001, N24770, AI669340, BF725049, AW298622, AI076755, AA489287, AI432423, AI885705, AI333169, AI828000, AA563808, AI139522, AI129404, AW069582, AA864279, AI192746, AW118992, AA939209, AA668916, AI564757, AW305369, AA847888, C75552, AA662165, AI696929, AI285881, AW606651, AU137938, AI814553, AA204900, AA056610, AW961767, BE255923, AI057234, BE746388, AI814745, C75086, AA599057, AA490113, AA834940, BF798693, AW028941, AA574248, AA897467, BG169516, AW407202, BF059031, BG030446, AA602132, AI921109, AI171602, BF336840, AI432792, F20465, AI268297, AA972138, AI121160, AA065277, AW815421, AW511846, AU157380, AW305193, AA065276, AA828531, AI049727, AA122378, BE241876, AA251814, N34210, AI567567, AA065275, AA971793, AI749941, AI221449, H69910, W25094, AW499564, C75461, N93237, R63270, BG170704, AA723814, AW293990, AL039399, AI240631, AA447796, H83038, AA598932, AW961429, AI094445, BE383532, BE669771, AW294261, BG012711, W57576, BF761772, AA131639, N80146, AA625946, BF855944, Z39516, AA441953, AA359095, H27740, BF921799, AW794926, AA378415, AA382847, AA701339, AI700223, BF743361, AA923716, AA328820, H69909, AI432486, AW977110, AW270206, AA809285, AA342845, AA339884, C00558, T10609, N44101, AI187829, AI340252, AI335635, M79208, AA765177, AA814298, R62825, AW590717, AW502811, AA204851, AA355746, AU155510, AA347918, AI470500, BG014393, AA709295, AA789162, AA862673, BE729337, R34037, AA251839, AA830387, AI185371, AW962476, AA314958, AW023841, AA625924, AW976799, AW015831, AW275618, AW502741, BF086375, AI828118, W03673, AI721111, R63222, R08239, T10371, C17620, AI361169, AA347917, AA171502, AA603411, AA095391, AA912518, W72108, R33496, AI868913, AA131721, AW518371, AA761154, N58100, BE746192, H95738, AV737030, AA382846, BE241628, T18550, BE925376, N87322, R45910, H82819, AI027852, AK026076, AK023418, AL137588, AB044750, AF263928, AK002050, AB044752, AK021850, AB044751, and AI498836.
HMSFT51	79	1091878	1 - 501	15 - 515	AV715134, AF139768, and AJ271684.
HAJCQ70	80	1091887	1 - 3267	15 - 3281	BE734515, AA446042, AA187765, AW963708, BE144726, AW388267, AA128252, BF238982, AA705646, AI745576, AA887587, AW269821, W79190, AI751939, AI860344,

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HAMFJ04	75	1091042	1 - 1659	15 - 1673	AV708059, BF207807, BE789361, AI768371, AI338520, BF128895, AI082511, AA250900, BE674806, AA491016, BE738988, BG026346, AI423838, AI915249, AI383087, BF955400, AA358276, AI567194, AW105110, BE745362, BF129513, AU117648, AC010186, AC006287, AJ271735, AL135910, AL441927, AL049561, AC008838, AC008952, AL121749, AC007160, U91323, AL139023, AL023579, AJ239322, AC007870, AC006167, AP000511, AL355305, AL121965, AL109843, AP001669, AC019155, AL138707, and AL109653.
HAIDK30	76	1091076	1 - 2177	15 - 2191	BG030257, BF662937, BG164999, BE467052, AW504354, BF508112, BE467049, AI888110, BE467051, AW962828, AI248937, AI084822, BG035687, AI249135, AA361076, AW469269, BF511869, AW674028, AA356078, AW296215, BE870263, AA897402, BE833830, AI161293, AI889765, AW885893, BF360992, AW804973, R22164, AF191545, and AC009073.
HCEIR41	77	1091358	1 - 728	15 - 742	AV719188, H67858, AW959062, D45273, AW960414, AW966380, AW961136, D80269, D59467, D80227, AW949500, AV726609, AW965153, AW966385, AW975613, AW966062, AV720791, AV702365, AA809122, AW959628, AW959597, AW965176, AW949586, D50979, AW973490, AV718489, AW975623, AW973326, AW966342, AV722801, D80378,

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HTEPC92	74	1091037	1 - 1649	15 - 1663	H02323, AI949415, D78845, AW966053, AW949642, AW975621, AV718489, D80166, AV719324, D59619, D80210, AW978634, AW949618, D80240, AW949643, AV723927, AV720211, D80212, AW949656, AW975618, D81030, D80219, AV724520, AW960553, AV719822, AV722801, AW966531, AV718692, AV718770, D80188, AW966534, AW966041, AW966062, AV720203, D51423, D51799, D58283, D80253, AW973307, AW949631, AV718844, AW949655, D59859, AV719557, AV720731, AW966013, AV699447, D80195, AV719783, AV718800, AV720464, D80391, AW959628, AW966054, AV719188, D80227, AV718440, AV720028, AW959570, D59889, AW965177, D80193, D80196, AV699550, AV699927, D59502, AW959202, D59927, AW973447, D50979, AW949653, D80269, AW978661, AW949629, AW949645, D80043, AW949657, AV699669, AW959597, AW975605, D80038, AV719468, C14014, D80366, D80022, AW966050, AW949654, AV721386, AV723097, AW965158, AW949641, AW949633, AW949632, AW966043, D80024, D59787, AW959582, C15076, AV700889, D57483, AV720812, C14429, AV718633, D80164, AV720791, D59275, D80045, AW960465, AW964488, AW949646, AW949658, AV699746, D59610, D50995, AV718931, AW959799, AW973485, C75259, D80378, T03269, D80241, D59467, AV720654, AW966022, AW964737, AV699682, AV645344, AV742001, AV742667, AW978648, AW973541, AV718707, AV701125, AV701335, AV701166, AV701043, AV701332, AV718938, AV701017, AV701248, AV742048, AV701431, AW966065, AV700229, AW965176, AV719049, AW959469, AW973334, AV645389, D80134, C14389, AW964756, AW966029, AW958993, AW965163, AW959136, AV719628, AV701419, AV742430, AV701154, AW973330, AV681510, AV681491, AW966075, AW958992, AV744770, AW959062, AW964477, AW956434, AW962245, AV701443, C14331, F13647, AW973474, AW752082, D51060, AV645343, AV699479, AW965197, AW956397, AW973488, D51250, AV701130, AV701149, AV701422, AA305409, AW960564, D80268, AV720878, AW965185, AW965184,

HOHEC71	69	1090538	1 - 616	15 - 630	AW971745, AFI52506, AFI52321, AB011160, AC005618, U88550, AR080280, and AB026436.
HCE3164	70	1090546	1 - 678	15 - 692	AA448371, AA448777, BE511209, AW341249, AFI92531, AB011176, and U27341.
HCE3L81	71	1090711	1 - 706	15 - 720	BF965765, AC005940, L42810, S83194, AB023658, and AFI117384.
HUFD162	72	1090753	1 - 972	15 - 986	AW175952, AU140869, T80660, AU137568, BFI10434, AC005626, AK023602, M22433, D90277, E03348, AC022516, E03349, D90278, L00692, L00693, and AC005794.
HE8FC96	73	1090865	1 - 3329	15 - 3343	AW957369, BF576270, BG181106, BE256459, AII25541, AW295443, AV718868, AV65054, BF981564, BE539459, AW957499, AI989947, BE568205, BE046999, AA779242, BE674435, AW236213, BF528673, AI167630, AW664585, AA836223, W69594, AA311403, N31338, AI272813, AI218709, AW296520, AA81816, AW182927, AI472964, AI367441, AA447971, AI934917, AA448115, AI075191, AI679378, AI199969, AI679887, AI355336, AW003822, N21384, AI808747, AI400127, AA004820, AI627481, BE672123, N26277, AA927143, BF028401, R91266, AA640518, BF700349, R67910, AW450415, AA761855, AI219952, AA455446, R91267, AA297775, AA373816, AI698552, W69408, BE245813, AW850061, AA860134, AA456858, AI813319, AI587528, AA350443, AW612171, BE925746, BF436580, AW975618, BE243697, AA830998, AW949645, AV724520, AW964468, AW966389, AW966330, AW949643, D80164, AW949642, C14331, AV718800, AV718692, AW960553, AV720791, D59502, AW966062, AV719783, AV720464, AW966054, AV719468, AW966053, AW949654, AW965175, AW964532, AW966013, C15076, AW965163, D59927, D80195, AW975621, AW960465, AW966050, AV699927, AW965185, AW965197, AW949646, AW949658, AA305409, AW966022, AW966534, D59275, AW966041, AV718489, AW973541, AW949631, AW965177, AW973330, AW966065, AV719822, AW978648, D59467, AV719557, D50979, AV719188, D80227, AW959597, AW959570, AW966029, AW959062, AW964477, AW949641, AW973334, AW966531, AW978634, AV718440, AV720028, D58283, D80022, D51799, D80166, AW959799, D80193, D59859, AW949655, AV699550, D59619, D80210, D80391, D80240, AW973474, D59787, AV719324, D51423, AW965184, AW973488, AW978661, AV720211, AV718844, AV718770, D81030, C14429, D80253, AW966075, AV720203, AW964756, AW973307, D80043, AV723927, AV718938, AW949653, AW949656, AV718633, AW959628, AW975605, AW949618, AW973485, AV718931, AV720731, AV699447, AW958992, AW958993, AV722801, AW959136, AW956434, AW964488, AW949629, AW962082, AW949633, AW949632, AW949657, D80269, AW959202, D80038, D80212, D80196, D80219, AW965158, AW966043, D80188, AV723097, AW975613, AW965196, AV718707, AW973482, C14389, AW960473, D57483, AW960454, AV720878, D80378, D80366, D80045, AW962245, D59889, D50995, D80024, AW966059, AV720812, AW956397, D59610, C14014, AW960564, AW753053, AW964737, AW959582, AF239798,

HE2PQ73	68	1089285	1 - 1246	15 - 1260	<p>AV717979, AV717953, AL036679, AV717986, AV701227, AV744771, AV718017, AL037601, AV717991, AW970679, AV745490, AV717995, AW975203, D80240, AL039504, AV741888, AV717996, AL039842, AV745366, AV718012, AX021518, AX020190, A85396, A25909, A85477, AR088705, A86792, A44171, AX042372, X68127, AX026824, AR062871, AR079804, AR037157, A84772, AR062872, AR062873, A84776, A84773, A84775, A84774, AX006821, AR067731, AR067732, A58522, A91750, AX035462, A20702, AX001082, A43189, A43188, A20700, AX026823, AX006825, AX006826, AX006822, AX046223, AR036905, AX003194, AR095491, AR095490, AR017907, AX001323, AX001324, AX035632, AX0035630, AX035631, AR072501, AR038762, A95051, A95117, AX026821, AX033488, AX033489, A38214, AR031374, A49700, AX001325, AX001326, A98767, AX033474, AR080470, AX033486, AR077142, I56772, I95540, AR018924, AX006820, A63067, A51047, A63064, AR018923, A48774, AX023549, A63072, A58521, A95052, A48775, AR068507, A18053, AR095492, AR068506, A93963, A93964, AX035629, A23334, A75888, I70384, AR043602, AR015960, AX006819, AX023550, AR043603, AR043601, AR000007, AR015961, AX023554, A18050, AR083151, A60111, AX023555, A23633, AX006817, A10361, AR020969, A23998, AX006818, AX023556, AR007512, AX009712, AX023552, AX023547, I60241, I60242, AX023551, AX006816, AX023546, AX023553, AX023548, I63120, I03343, AX012337, AX003207, AR054109, AX008555, AR031375, AX006823, AR022240, A58524, A58523, A81878, A24783, A24782, AR069650, E12615, AX018504, AR035193, A92133, E14304, AR096518, A27396, AX000085, AR027100, I28266, AX042377, AR091518, AX042375, AR085082, AR085089, AR085091, AX042373, A49045, AR085079, E16678, A82653, AR085083, E16636, I06859, A93016, AR093392, I25027, AX033490, AR069426, AR000006, I26929, E13740, AX006824, I44515, I26928, I26930, I26927, A58525, AX033487, AX009487, AR069374, I49890, AR069375, I44516, AK026727, A58526, A91753, AX002551, AJ244003, AR072503, AR072500, AR025207, AR072502, I18371, AF261960, AF261963, AF261959, AF261962, AR080280, E06034, AF156296, A13038, AX027488, A29289, AR029417, AR088315, AF156294, AR067733, AR029418, AR067734, AX025617, A64081, AR028669, AR028668, AR028667, AR028670, AR017908, A68112, A68104, A98467, I62368, AR031488, I13521, I52048, AX047291, I44531, A84746, AR093383, A17115, AR028672, A18079, AX012320, AX012321, AR038066, AR074025, AX047314, AX047311, I15353, I66495, I66494, I66498, I66497, I66496, I66486, I66487, A71440, A71435, AB037923, I13349, A07699, E08322, I74623, AF261964, AX020192, AF261966, A60109, AC018903, AC068470, AL049186, AL049186, AL022167, and AC022305.</p>
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HE8PB92	66	1086828	1 - 685	15 - 699	N29316, BE892536, AW383428, AW403946, N30453, W03476, AW407871, N35519, W02963, N20533, AW403434, BF896859, BF794609, AA837208, W02298, BE559773, N46511, AI439520, BE513826, N29315, BE384088, BF572179, BF303959, BF572042, BF690655, BF203242, BF692515, BE275832, BF305905, AW402824, N28949, AW402928, and BE729612.
HTMU66	67	1086856	1 - 1077	15 - 1091	T68139, AW937136, and BF808052. AL039924, AV731085, AV738934, AV718844, AV718018, AV737584, AV745350, AV718002, AV717989, AV701782, AV717988, AV742995, AL045794, AV745723, AV718016, AV745724, AV743601, AV701012, AW013814, AV735727, AV746102, AV758878, AL036630, BF508972, AV700229, BF294063, AV745917, T02921, AV719000, AV717956, AV699447, AV717963, AV701017, AV741012, AL044412, AV717962, AV717983, T24119, AV717984, AV717990, AV745488, AL044364, AV745080, AV720464, T24112, AV719568, AV722801, AV701248, AV717980, AV701043, AV718023, AV718021, AV717966, AV746162, AV717959, AV701118, AV746335, AV701166, AV717965, AV723927, AV717946, AW450335, BE439760, AV717972, AV742667, AV701163, AV742720, AV718858, AV718681, AV720220, AV718010, AV717970, AV742001, AV717993, AV736165, AV701154, AV744934, AV745369, AV743654, AV743008, AV720731, AV717976, AV701261, AV718003, AV718001, AL039476, AV717978, AV742862, AL039386, AV740535, AV720607, AL045341, AV717971, AV724520, AL040992, AL039109, AL038531, AL037726, AL039629, AL039625, AL039648, AL038837, AL039074, AL039678, AL039108, AL039538, AL039564, AL039156, D51250, AL039659, AL039566, AL039509, AV745392, AL039521, H00069, AL039128, AL044407, AV745831, AL036973, AL045337, AL037051, AL045353, AL039423, D59787, AL042909, D80253, AL039410, AV700889, AV717949, AL039150, AV717960, AL044530, AL038821, AL038025, AV718020, AL036725, AL043422, AV717941, AV701125, AL043445, AV718008, AL039459, AV718011, AV717948, AV701431, D80219, AW976625, AV717985, AV701332, D59275, AL043586, AL043423, AV699479, AV718692, AV717968, AW973447, AV699927, AV721784, AV717973, AV718489, AV717987, AW064110, AW969383, AL043441, AV717952, AV701428, AV719783, AV720034, AV717967, AV700622, AV701121, T23947, AV717943, AV701004, AV717964, AL036650, AV744770, AL036196, D51423, AV717955, AV717992, AL037639, AV717974, D80043, AW451070, D59619, AV718014, AL037615, AV744773, AV718006, AL535983, AV718013, AV717958, AJ293456, AL036767, AV701335, AV701443, AW969322, AV717997, AW975312, AL039085, AL535783, AL036117, AV746385, D80227, AV717957, AV701149, AV735369, AW063533, AL037526, AV718000, AV717961, AL042334, AV701357, AV744768, AW452756, AV720812, AV717954, R47228, AV699669, AL036238, D80366, AV717942, AV718007,

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HDQK06	61	1084465	1 - 2622	15 - 2636	BF899863, BF330510, BE410120, BE388453, BE388922, BE389727, BE389752, BE390985, BE389948, AW971745, BE389145, BE386347, BE613349, BE612945, BE622828, BE613070, BE613136, BE622931, BE621911, BE622868, BE614230, BE614307, BE614237, BE618811, BE622925, AI083689, BE620742, BE614451, T05706, BE620756, Z99396, AI433297, AW861889, AW858455, AW363220, AW804686, AW392670, AW384394, AW861944, BE695785, AW604726, AW604723, AW372827, AW858526, AW858525, BE705903, BE705906, AW577135, AL119497, AL119319, U46350, U46347, AF134707, AF019887, D50410, A61271, A61272, AR080280, I25027, AR054109, I44515, I26928, I26930, I26927, I25041, AR069374, AR069375, I44516, AR091518, AX009487, AR093384, AR035224, I85513, AR009152, AR009151, AR027099, AR093392, AR093383, AR019098, A94046, A94054, AR020199, AR020200, AR001287, AR020198, AR020197, I05393, A10617, AR028792, AR029418, A01324, A01323, AR019094, AR034783, AX030966, A92666, AR067733, AX009486, A92668, AX029455, I49890, A92667, AR067734, AR067731, AR064322, AR064323, AR064320, AR064321, AR067732, AR029417, A83642, A94048, A94061, A83643, AX035462, A92665, AX027785, AR038307, AR038321, A49045, I63120, A70359, I89986, A32110, AR083151, AX003194, AR028791, AR028793, AX024906, I05430, I09121, AX032758, AB026436, AR055065, AX046357, A30600, AR055066, A81671, AR060234, AJ251859, AX030435, AR054110, AJ279014, AR066494, and AR069079.
HNGME19	62	1084997	1 - 1425	15 - 1439	AI039143, AA625076, R91027, AW956093, R88767, AW956097, T04933, AA092577, BF382317, N83651, AF169693, and AK025871.
HELGO48	63	1085552	1 - 1611	15 - 1625	BG122833, AI088936, AI681970, AW475066, AW517258, AA447640, AW067824, AI432519, AI089690, AI969201, AA076638, AI362134, AW026926, AA193257, BF769770, AA916592, BG006202, AA377530, F25044, AA305704, AA303642, AA448044, AI471047, BE926036, AI739091, AI014970, F35714, AA079606, AW067881, AF060567, AL391688, and AL110206.
HRDAF83	64	1085723	1 - 1006	15 - 1020	BF749563, AI825217, AW604115, BG258877, AA316377, BG013602, AF109681, and AF137378.
HBJHS02	65	1085743	1 - 733	15 - 747	BE560978, AW402242, AW402801, AW383418, AW383426, AW383396, BE271199.

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HTLDS55	60	1083991	I - 1942	15 - 1956	<p>W92236, AA299662, AA004847, BE149252, H56718, T77776, AA002009, AA227236, AI922495, AA722941, AA456022, AA299663, AA001788, AW630463, H56641, AW904581, AV733999, AW971745, AL119457, AW392670, Z99396, AW804686, AL119319, AW861944, BE695785, AW604723, AW858526, AW858525, AL119355, AL119324, AL119497, AW877209, U46350, U46351, AL119363, U46349, BE705903, BE705906, AL119391, AW577135, AW372827, AL119483, AW384394, AW861889, AL119341, AW858455, AW363220, U46347, BF868687, AL119484, BE119443, U46341, AL119444, BF868697, U46346, AW604726, AL119439, BF868684, BE705905, AL119522, AI142134, AL119396, AL119335, AL043033, AL119401, AL134538, AL134542, AL134528, AL134902, AL134531, AL134533, AL119418, BE705904, AL119399, AL042984, AL119496, AI142132, AL134525, AL134536, AW861954, U46345, AL119464, AL042450, AL042614, AL043029, AL042544, AL043011, AL043019, AL042542, AL042965, AL042975, AL043003, AL042551, AL132826, AB040902, AF169677, U42975, AB026436, AR066494, AJ251859, AX030435, AR060234, AR054110, A81671, AX046357, AJ279014, AR080280, and AR069079.</p> <p>AI890919, BF222811, AI018797, AA913452, AI797580, AV709407, AV726067, AV726505, AV702798, AV707783, AV659189, AV706183, AV701611, AV728270, AV702427, AI809012, AV704592, AV704974, AV706724, AV702637, AV707882, AV658362, AV725369, AV697638, AV704916, AV727932, AV728777, AV729983, AV706910, AV725927, AV728872, AV707769, AV726738, AV706746, AV705020, AV704116, AV732002, AV727189, AV728289, AV706851, AV724987, AV701538, AV704611, AV727103, AV707088, AV726480, AV728715, AV706047, AV703591, AV702869, AV723449, AV647654, AV656240, AV701728, AV703417, AV731759, AV652156, AV702671, AV728455, AV703388, AV707948, AV705504, AV702787, AV726392, AV730781, AV706527, AV745906, AV702984, AV732149, AV703232, AV751921, AV702026, AV709935, AV702954, AV752043, AV753374, AV730288, AV707589, AV704924, AV707117, AV701237, AV731915, AV702409, AV662191, AV707798, AV702537, AV709356, AV725281, AV705234, AV729129, AV701320, AV652547, AV707171, AV704279, AV705263, AV731744, AV731313, AV703012, AV726520, AV707686, AV751555, AV731977, AV726830, AV729357, AV731043, AV701783, AV727355, AV705047, AV706104, AV685688, AV652808, AV702498, AV706453, AV707690, AV731694, AV732353, AV645778, AV730711, AV730165, AV709897, AV732155, AV646736, AV731708, AV731275, AV725387, AV706889, AV752443, AV706220, AV727576, AV726559, AV729220, AV707685, AV702958, AV725152, AV703367, AV761002, AV730547, AV732653, AV730171, AV732746, AV708347, AV762691, AV706025, AV705416, AV706234, AV701496, AV706989, AV650367,</p>
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4					<p>AI821714, AC008812, AP001753, AL138726, AC005098, AP002085, AC020908, D83989, D84394, AL024509, AL133445, AC004867, AC005540, AC007957, AP000556, AL031311, AC004826, AC004638, AL133284, AF038458, AC002082, AC011475, AC025594, AC008474, AL096701, AP000557, AC005523, AC009086, AL034406, AC002430, AP001748, AC024082, AC002558, AC009123, AL133241, AL009181, AC007384, AG002546, AC008623, AL162615, AF190464, AL163249, AC005962, AC016903, AC009032, AC008616, AF053356, AC010489, AC004686, AC011510, AC010422, AC010553, AL121658, AC004966, AC005231, AL354872, AL034380, AC005632, AC005365, AC008403, AL121928, U78027, AC005215, AP001630, AL049795, AC008567, AC002470, AC007371, Z93015, AL035422, AL157688, AC008760, AP001412, AC006038, AF126531, AL049757, AC006483, AC018720, AC004134, AC024561, AC006367, AL022147, AP000152, AC007850, AL022322, AC083863, Z94801, AC005519, AC004668, AC008537, AL109623, AC003007, AL163279, AL163282, AP001717, AF135026, AJ003147, AC002316, AP001053, AL121845, AC011444, AC007842, AC005291, AC005250, AC001231, AF168787, AC011469, AL160175, AC007292, AC005840, AL121943, AL023575, AL080317, AF165926, AC006251, AC004858, Z49237, AC006111, AC023490, AC003029, AL356652, AL137881, AC006009, AC006285, and AL121601.</p>
HOFMN17	57	1083062	1 - 778	15 - 792	<p>BE781884, BE255612, BF842043, AA683020, BE293969, BF841112, BF303486, BF930132, BF771921, BG251102, AW409699, BF812536, BF923844, AI809532, AI453056, BF744747, AA778576, BE254325, BE041585, AW005786, AI042586, AI359703, H91861, AW205217, H91914, AF307079, AB020628, AF111097, AF111098, U72487, AF081144, AF081146, U78105, AF111099, AF081145, and AF081147.</p>
HKABY22	58	1083735	1 - 1062	15 - 1076	<p>BF317466, BF737694, AW971745, AW604723, AW392670, AW804686, AL119324, AW858526, AW858525, BE695785, AW861944, AL119497, AW577135, AW372827, U46347, AW384394, AW861889, BE705903, AW858455, AW363220, BE705906, AL119363, AL119457, Z99396, AL119443, BE705905, AL119319, U46341, AL119396, AW877209, BF868697, AL119484, AL119341, AL119391, AL119355, U46346, U46350, AW604726, U46351, U46349, BE705904, AL119335, AL119444, AL119483, AW861954, AL119496, AL119399, BF868687, AL134533, AL134528, AL119522, AL119439, BF868684, AL134531, AI142134, AL037205, U46345, AL134529, AL042614, AI142132, AL134527, AL134538, AL119418, AL043003, AL042450, AL042965, AL042975, AL042542, AL042544, AL042970, AL043019, AL042984, AL043029, AL119511, AL042551, AL119488, AL119464, AK025342, AL137477, AF217289, AB026436, AR080280, AI279014, AI251859, AX030435, AR054110, A81671, AR060234, AX046357, AR066494, and AR069079.</p>
HLDOJ74	59	1083901	1 - 2373	15 - 2387	<p>AW959618, AL528457, BE857709, AL528430, AI806250, BF126180, AA455382, AI084580, AW368035, AA005065, AL529481, AI088155, AI566044, W92235, AA706063, BE328495,</p>

					<p>AW965185, AW965197, AV718707, AW949618, AW964737, C15076, D57483, D80164, AV700889, AV718931, AV699669, AW966022, AW959799, AV699746, AW978648, AW966065, AW966043, D50995, AV720654, D59610, C14331, T03269, AW965175, AW966075, AW973541, AW966029, D59787, D80378, C75259, D59467, AW973334, AV700229, AV718938, AW959062, AW964477, AW959469, D80241, AW958993, AV719913, D51060, AW964756, AW973474, AV742001, AV742667, AV699682, AW965163, AV701125, AV701335, AV701166, AV701043, AV701332, AV701017, AV701248, AV701431, AW965184, AW973488, C14014, AW973482, AW958992, AW959136, AW956434, AW962082, AW962245, AV719628, AW975613, AV742048, AW752082, AV681510, AV681491, AW965196, D81026, AV719049, AW960473, AA305409, AV720878, AV701130, AV701419, AV701154, AW973330, AV645389, AV645344, AW956397, AV718530, D80134, AW965176, AW966059, AV701443, AV701004, AV742430, AW960364, AW960454, AW960532, AW966023, AV699479, AW753053, AV645343, AV681468, AV701149, AV701422, F13647, AW960504, D80268, AW966030, AW960414, AW973465, AV681529, AW178893, AW975623, D58253, AV720220, AV701428, AW966560, AV701415, AV701344, D80949, AW960570, AV719000, D51250, AW973470, D80168, AV701357, AV745080, AW966032, AV700622, AV719727, D81111, D51022, AW179328, AW960474, AV721784, AW177440, AV744690, AA305578, AV700159, AW949630, AV720150, AW966049, AV741012, AV699652, AV720607, AV720533, AV701021, AV744934, C14227, AV724977, D59695, AW964532, AX039409, AX039410, AX039412, AB046859, AX027925, AR070327, A62300, A62298, AX033851, A84916, AX047063, AJ132110, AX047064, X67155, A78862, AX020191, A67220, AX020190, AR018138, AX021518, A25909, D89785, D26022, AX047062, Y17188, D34614, AJ302649, AX035434, AR092424, D88547, AR025207, AF058696, AR008278, AB028859, AR077702, AR087649, X82626, AX028130, AJ294956, AF260572, Y12724, X68127, AX015396, AB012117, A82595, AR088705, AR060385, AB002449, AR074545, A85396, AR074141, AR066482, AX042372, A85477, AJ287395, A94995, I19525, A86792, A44171, AR008443, X93549, AR016808, I50126, I50132, I50128, I50133, AR066488, AR016514, AF135125, AR060138, A45456, A26615, AR052274, I14842, AR074139, Y09669, A43192, A43190, AR038669, AR066487, AR066490, AR074136, A30438, I18367, D88507, AR054175, D50010, Y17187, A63261, AX035429, AX035428, AX035426, AR064240, AB037923, AR008277, AR008281, AR008408, AR091537, AR062872, A70867, AR093385, I79511, AR016691, AR016690, U46128, A64136, A68321, AB033111, D13509, AR060133, U87247, AR087528, U79457, AF123263, AX014811, AR032065, X93535, and AR008382.</p>
HMMCHO	56	1082413	1 - 766	15 - 780	<p>AI049671, AI034206, AW079877, AI791913, BF741641, BF681546, AI792133, AU144517,</p>

HPLBQ44	54	1081779	1 - 2667	15 - 2681	AR032065, X93535, and AR008382. AL519790, BF795638, BF026182, AW994029, BE394273, BF205114, BF027356, BE895131, BE872776, AL519789, BE410894, BE276329, BE276487, AW957340, BE388047, BE391321, BE394498, BE743519, BE384608, BE408222, BE409787, BF689435, AL043880, AW972147, BE269716, BE311514, BF027650, BE646542, BE617138, BE728198, BE388823, BF593614, BF736579, AW005102, BE019544, BF125559, BE542482, BE905771, BE894272, AA137033, BF155204, BG010268, AA810411, AA523117, AI671452, BE385576, AI339682, BE206310, BE673474, AA437080, BE871373, BG121363, W90768, AI091057, AW606873, AI288535, BE388630, AA528033, AA137115, BF798691, AI338926, BF765478, AW130160, AI683304, AW083986, AI859189, AI199890, AI199896, AI625514, AI269181, AW194157, AA114896, AA114897, AA922395, AI990810, AI349372, AI168791, AI812097, AI274942, AA128536, AI266219, AI636417, W90701, BE933377, AI862000, BF847189, AW204921, AI743815, AI023431, R51136, AI091111, BF002998, N78924, AI28362, H48319, AI305176, AI873659, AI056132, AI087400, H48227, R52200, R62400, AW339362, H79689, AI886641, R62399, AI680503, BF822055, F11141, AA938005, AA423981, BF822056, AW583151, BE727538, AI350674, M78260, W05294, AA330985, Z46207, BF904232, BG009686, AI915189, BE185674, BF829979, BE185861, R12869, BE714832, BF762421, AA523876, BF847074, BF758304, BE829339, R38443, BE294165, F08308, F05074, AA339020, BF953251, R51028, H79690, BF081236, BE770844, F08811, AI982566, AA343945, AW947392, F04534, R43758, BE185748, AA368664, BF989075, BF988558, AI801276, AW089229, BE185979, D78779, BE671521, BE939678, AA465535, AA368064, BE706060, AI751158, BF995582, BF987557, BF084981, BE698927, AW062626, AW904903, T05279, and AK025157.
HAHBE26	55	1082286	1 - 862	15 - 876	AW862431, AW975621, AW966053, AV719783, AV720464, AW975618, AV719468, D80193, D59502, AV723927, AV719188, D80166, AW960465, AV719324, D80195, AW966013, AV720211, AV718800, AW949641, AV724520, AV720203, AW966531, AW978634, D59619, D80210, D80240, D80212, AV718770, D51799, AW966054, D81030, AV718844, D80227, AW966062, AW959582, D58283, AW959570, AW949656, AW949642, AV699927, AV719822, D80219, AW966534, D80391, AW949658, AV718692, D51423, D80253, D50979, AV718489, AW973307, D59859, D80269, AV720028, AW959628, AW949646, AV719557, AV720731, AV699447, AV722801, D59889, AV699550, D80038, D80196, AW966050, AW966041, C14429, D80188, AW960553, AW949653, D59927, AW949631, AW949643, AV718440, AW965158, D80043, AW959202, AW949657, AW973447, AW959597, AV723097, AV720812, D59275, D80022, AW949632, AV720791, AW965177, AW949629, D80045, AW949645, AW949633, AW949654, AW978661, C14389, AW949655, AV718633, D80366, AW973485, AV721386, AW975605, D80024, AW964488,

AV720791, AV721386, AW966050, AW949632, AW949654, AW973447, D59275, AW949653, AV700889, D80022, AV720812, AW966043, D80024, AV723097, C75259, AW949646, AW949658, AV718633, AW964488, AA252405, AW959799, D59502, AW960465, D80378, D59610, D80045, BE245697, AV718931, T03269, AW964737, AV699669, AW973541, C14014, D50995, AV720654, AW973485, AW966022, AV699682, D80241, AV645389, AV742001, AV742667, AV701125, AV701335, AV701166, AV701043, AV701332, AV701017, AV701248, AV742048, AW958993, AV701431, AW965176, AV700229, C15076, AV645344, AW978648, D80134, AV718707, D59787, AV718938, C14331, AV699746, AV681510, AV681491, AW973334, D80164, AW959469, AV719049, AV701419, AV742430, AV701154, AV745080, AW964756, AV719628, AW959136, AW966029, AV719913, AV720220, AV699479, AV701443, AW752082, AW965163, AV645343, AW958992, C14389, F13647, AW959062, AW964477, AW956434, AW975613, AW973474, AW966065, D51250, AV701130, AV719000, AV701149, AV701422, D80268, D51079, AW956397, D58253, AW965197, D59467, AV718530, D81026, AW962245, D51060, AW966560, AV681529, AA305409, AW965185, AW965184, AW973488, AV720878, AV701004, AW962082, AV701428, AV701415, AV701344, AW960414, AW973330, AW966075, AW973470, AV720607, AW965175, AW178893, AW973482, D80949, AW975623, AW973465, AV681468, AW960504, AW960564, AV721784, AW960473, AW966059, AV718674, AW965196, AV699411, AV741220, D81111, AV700622, AW753053, D80168, AW966023, AV742732, AV744770, AW960454, AV701021, AW960570, AV741221, C14227, AV724977, AV719727, AV645383, AV645393, AV681528, AV681474, AV645339, AV681472, AV681507, AV681465, AV681525, AC006252, AF259262, AF259263, AF245704, AB045180, A62298, AR070327, A62300, A84916, AX033851, AX027925, AJ132110, AX047063, AX047064, X67155, AX020191, A67220, D89785, A78862, AX020190, AX021518, A25909, AX035434, AR018138, AX047062, D26022, D34614, AJ302649, Y17188, D88547, AR025207, AR092424, X82626, AR008278, AJ294956, AF058696, X68127, AF260572, AB028859, AX028130, AX015396, AR077702, AB012117, AJ287395, AR087649, AR088705, A85396, AR074141, AR066482, Y12724, A85477, X93549, A44171, AR074545, I19525, A86792, A82595, AX042372, AB002449, A94995, AF135125, AR060385, AR008443, I50126, I50132, I50128, I50133, AX035429, AX035426, AR066488, AR016514, AR060138, A45456, A26615, AR052274, AR064240, D88507, AB037923, AR074139, AR054175, Y09669, A43192, A43190, AR038669, AR008277, AR008281, AB033111, A30438, AR066487, AR074136, Y17187, AR066490, I14842, I18367, D50010, A63261, X64588, AR016691, AR016690, U46128, AR008408, AR091537, AR062872, A70867, AR093385, Z32749, D13509, A64136, A68321, U79457, AR060133, AR087528, I79511, U87247, S69292, AB023656, AF123263,					
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HMTAK58	50	1071812	1 - 659	15 - 673	and AR069079, BF103655, A718659, A1651912, BG232105, AW956578, BE049585, BE247532, AA743043, BF513581, AA029932, AW408031, AA028937, AA380908, AI302421, AI984467, AI926944, BG231836, AW268886, BE675569, AW512035, A1635345, AA884484, BE247537, N67007, BE929465, BF509920, AA506046, BG236197, BE244322, AI873194, AA748575, R00578, AA749217, AI633684, AA748305, AW754406, BF895003, AA815159, BF895254, AF133299, AF285087, AF285089, AC007068, AF285088, and AC010186.
HPWBS63	51	1073467	1 - 722	15 - 736	BF900226, AR075074, and AR075073.
HPWBS63	52	1077575	1 - 820	15 - 834	AW024340, BG152772, AW024329, AI885035, AI870591, BE857493, AA644167, BF956587, AI819124, AI334166, BF998808, AI017361, AI954925, AI140418, AA890034, AI242827, AA805756, AI921497, AI401584, AW439173, AI359168, AI096461, AI027350, AI359373, AI568727, AI052420, AA262805, AA037812, AI435579, AA873297, AA037813, AA516397, H38962, AA830180, H03939, AA830569, H21198, H03127, AA055952, H11109, BE646587, AA830181, AI091595, AW058142, AI143045, AI819814, W93685, H20951, AI026037, AA846280, H41747, AI383728, W93737, AA843752, H21839, AI566942, BF149291, BF094494, BE719277, AI934654, H41880, BE932659, BE933275, BE933281, AA643549, BE701088, H11207, BF088159, H41748, AI159926, BF954997, BE719283, W84562, AA769546, AA055953, R77647, BF813408, AI797820, AI094168, BE932667, AI928723, BF002144, AI202826, AI276117, AI283864, BF316030, BE720028, AI910381, BF062494, W84485, BF752662, BE932599, BE719290, AI638337, BE932592, AI652400, AI564949, BE719284, AI885162, AA091517, BE719313, BF476370, AA059093, BF726765, AI201513, BF803498, BF820891, AW364108, H30025, BF840522, BF804647, BF803499, BF804575, AA261929, BE719267, AW197525, AA059092, BE696429, BE696430, AI422045, AI356155, AA677506, AI589762, BE933280, AA743504, AW903400, BE762317, BE762345, BE144808, AB020718, AI289016, AI289017, AK000605, and AW243552.
HSDJH63	53	1078285	1 - 1500	15 - 1514	AA741118, AW451491, AA252360, AW975618, AV718489, AW966053, AW978634, AW960553, D80253, D80188, AV722801, AV699447, D51423, AV724520, AW966531, AW966534, D58283, AV719557, D59859, AV699550, D59889, D80166, AW978661, D59619, D80210, D51799, D80240, AV719822, AW949645, AV718692, AW973307, AV719324, AW965177, AV720211, AV720731, AV723927, D80212, AW949656, D80227, D57483, AW949642, D80366, D80195, AW975621, AW959570, D80219, D81030, AV718440, D80391, AV720028, AW966054, AV720203, AV719188, AW966062, AV719783, AW966041, AW959628, AV718800, AV718844, AV720464, AV718770, AW949657, D80269, AV699927, AW949618, AW966013, AW959202, D80196, AW949641, AW959597, D59927, C14429, D80038, AW949629, D80043, AW949631, AW949643, AW949655, D80193, AW965158, D50979, AW959582, AW975605, AV719468, AW949633.

HNFHW14 HUKDS43	45	1057738	1 - 387	15 - 401	BF183209.
	46	1061241	1 - 1691	15 - 1705	BE271587, BF437289, BE879350, BF446399, BE736154, AW991607, BE813144, AW373239, AW955378, AW372628, R73350, BF842394, BG248958, R51402, N27996, R54681, BF570406, AW749924, AA377857, AA422157, AW991461, R50597, AI808730, BF570043, AW991517, AW393029, AA326416, AW818812, BF939464, BE548597, AW373220, BE513102, AI827898, AI825876, AI650385, AI827701, AI888306, AV709323, AI934499, AW577304, BE513101, AW006103, AI422225, AA524283, AI088893, AI422224, BG251334, AI217369, AI380811, AI469281, AA494534, AA975272, N21338, BE513513, AI359575, AX033274, and AX014703.
HAOAD02	47	1063218	1 - 5092	15 - 5106	BE349027, AA460958, AA460959, AI291926, AA461266, N58870, N72734, AI277327, AA461265, H43656, BF927468, H44722, BE835623, BE835624, Z33537, AI903814, BF366858, AA430219, AW968357, AA485068, AA463480, AF112345, AF074015, and AF172723.
HOUCD83	48	1070724	1 - 3878	15 - 3892	AL043329, BF965834, BG167996, AI740473, AI970994, AI741566, AI817205, BF034229, BF037696, BE644852, BF540943, AL529179, BF439276, AI740889, BF678309, AI743775, BE464341, BF801570, BE544927, AW466917, AI435153, AI091411, AI091191, BG026179, AA129716, AW117751, AI274064, BG236187, BF439185, AA643517, AI952321, AI675270, AI753250, AI142000, AI436790, AA133238, AI220185, AW204898, AW468944, AA421937, H01552, R79134, AA824389, AI819042, R78584, R78832, R69250, R78848, AW303628, R66023, Z39338, BF245312, BF085261, R74298, AI309540, AA970121, R67482, AV728867, AI742992, T40950, AI886090, BF980298, R68981, AA406320, R32257, R31056, R24118, BF526590, AA905273, AW182341, R33781, R66024, R24066, R40089, R32835, R73089, R68952, R62781, R69132, T40020, R68982, H01458, BF054965, R33666, R79337, R73088, R79327, R31541, BE171967, R79038, N55916, R62780, R32938, AL050071, AF282874, AF195833, AF195834, and AF195835.
HOFMS43	49	1071027	1 - 580	15 - 594	AW971745, AW392670, AW804686, AW604723, BE695785, AW861944, AW372827, AW858526, AW858525, AW384394, AW861889, AW858455, AW363220, AL119497, BE705903, BE705906, AW577135, U46347, U46341, AL119443, AL134531, AL119457, AL119319, AL119324, AW877209, Z99396, BE705905, AL119396, AL119484, AL119363, AL119341, AL119391, AL119355, U46350, AW604726, U46351, U46349, AL134528, BF868697, BF868687, AL119444, AL119483, U46346, AL119335, AL134920, AL134524, BE705904, AL119399, AL134533, AL119496, AL119439, BF868684, AL119522, AW861954, AL037205, U46345, AL043029, AL042614, AL042975, AL134538, AL119418, AL042965, AL042450, AL043003, AL042542, AL042544, AL042970, AL043019, AL042984, AI142134, AL119511, AL042551, AL119464, AL119488, AB026436, AR080280, AJ251859, AX030435, AJ279014, AR054110, A81671, AR060234, AX046357, AR066494,

					AW979176, AW969885, AW973219, AW975628, AW974802, AW975632, AW975971, AW979169, AW973750, AW975027, AW974823, AW976982, AW975981, AW970969, AW979106, AW975990, AW975020, AW970942, AW974975, AW969643, AW975254, AW972849, AW976511, AW975649, AL119399, AW974785, AW973185, AW969852, AW975692, AW979294, AW971968, AW970029, AW969703, AW972296, AW975585, AW972817, AW974806, AW972867, AW979220, AW979142, AW979219, AW970010, AW971378, AW971326, AW976000, AW973130, AW974338, AW976506, AW972857, AW970962, AW974792, AW974971, AW970936, AW973270, AW970113, AW971404, AW972884, AW973202, AW973805, AW972719, AW975244, AW976515, AW975976, AW979165, AW971387, AW976510, AW973770, AW975975, AW969884, AW979054, AW972943, AW969759, AW979083, BF592735, AW970587, AW973650, AW979232, AW979175, AW973986, AW979064, AW975938, AW975138, AW975968, AW970921, AW969766, AW979116, AW972806, AW973987, AW975910, AW975943, AW973819, AW973164, AW970097, AW973824, AW971964, BF588769, AW975904, AW974379, AW969752, AW973967, AW975933, AW973207, AW972695, AW972882, AW973104, AW969782, AW970868, AW973985, AW973210, AW971129, AW973209, AW972440, AW979178, AW972883, AW979081, AW973167, AW975162, AW972637, AW970110, AR035695, AF068288, AB013102, AC026884, AR060234, AX046357, AR066494, AJ279014, A81671, AR023813, AR064707, AR069079, AX030435, AJ251859, AR096545, AB026436, AR054110, and AR096546.
HTENW06	41	1040030	1 - 1131	15 - 1145	BF724829, AA311043, H11578, AW665008, R59755, AA927842, AA199784, AI000363, AI126309, R13187, AI693684, and AL096806.
HODFW19	42	1052332	1 - 1374	15 - 1388	AL119917, AL138095, AL119463, BF511055, AI739217, AI805761, AI870737, AW973926, AL526882, BF924366, U83567, BG252755, AI075929, BF948710, AC005618, AF152513, AF152326, AF152510, AF152509, AF152323, AF152318, AF152507, AF152508, AF152322, AF152503, AF152319, AF152506, AB011160, AF152321, AB002325, AR025269, AF152514, AF152515, AF152325, AF152512, AF152327, AF152324, AF152511, AF152328, AF152320, AF152505, AF152516, AF152329, AF152504, AF152517, AF152530, AF152518, AF152331, AF152495, AF152493, AF152500, AF131761, AF152489, AF152498, AL117449, AF217749, AF152502, AC005754, AC008688, AF152499, AC025436, AF152497, AC005752, AF152494, AF152491, AR025266, AC074130, AF152492, AF152527, AK023190, AK021915, AF217757, AF152490, AF282973, AF152501, AB046841, AF152496, AF152528, and AR025241.
HACAD35	43	1052372	1 - 598	15 - 612	BE743922, BF037138, BF727228, BF726450, AA346825, AA228126, AC007363, and AC007363.
HHPDX35	44	1056108	1 - 751	15 - 765	BF038374, BE061260, BF379151, BF379150, BF351968, F24300, and AW376415.

HPMSA16	37	1011769	1 - 1036	15 - 1050	BG110380, AI671613, AA075344, BE550608, AA778899, AA496875, N38985, AI888310, AI088077, AI061591, AI493142, AI338275, BE075101, AW872997, AA075597, AA136180, AI459760, BE645791, AA909786, AI292044, AA279124, AI871819, AA369669, BF912395, AA136299, N38971, AI000293, AI924804, AA952968, AW022179, BE910012, BF909523, H09755, AI244527, AI114660, AI003708, BE167338, N48131, BF908708, AA279243, Z39538, F04624, AW384456, R22931, T19719, N48117, R23036, AW384458, AA279928, AW386819, AW842474, T19720, H47088, AL036212, AA157797, AW752757, AW376997, AW376921, AV734931, AI933835, AB029028, AF180425, AF114491, AK024454, AK024426, and AC004262.
HITKE35	38	1027025	1 - 1495	15 - 1509	BE677799, BF432054, BF061402, AA922217, AW451106, AI949280, BE218951, AW451993, AA454675, N35987, R26762, AA456121, R25954, AW470239, AA204920, AI669723, BE855992, C02224, and BF061357.
HCRPE25	39	1034245	1 - 1177	15 - 1191	BF056859, AI950360, BG236479, AA418443, BF591089, BF475316, AA234138, AI128275, AI949635, BF224126, AA236217, N31643, AI188960, BF224128, N27174, AI681942, BF447701, N39919, BF594184, AI681350, AA557621, N57105, AW293218, AI703156, AI765582, N31647, AI735524, AA172040, N30714, AI767963, AA172194, AV713103, H93692, BF115111, AW105367, AI000265, AI033224, N41328, N57098, R08399, AA418393, AI474441, AA125975, BG230833, AI767420, AR035695, AF068288, and AB013102.
HWAGX4 9	40	1034246	1 - 1669	15 - 1683	BF056859, AI950360, BG236479, AA236217, AA418443, AI361031, BF108683, BF591089, AA234138, AI128844, BF475316, BF594184, BE670221, AA757278, AI342632, AI361934, AW293218, AI765582, AI703156, AI128275, AI949635, AA056658, BF224126, AW057598, AI361461, AA056410, AI830753, AA172040, AI146443, AI767963, N31643, BF224128, N27174, AI829376, AI188960, AI949283, AI335683, AI681942, AI290703, AA172194, BF447701, AA557621, AI681350, AA128162, N31647, AW237789, AI033224, BF589027, AW004907, AI735524, AW515103, AW305109, N30714, R62698, BG153411, R76068, AV713103, H93692, AA125975, BF115111, AW105367, AI000265, R33815, AW853796, AI474441, N39919, AI770139, AA723866, R08355, N41328, R63659, R33814, N57105, R75891, AA703473, AW016556, AI700946, AA418393, R08399, AW510566, AI767420, N57098, BG230833, AW979127, AW972292, AW971975, AW974101, Z99396, AW975037, AW976023, AW975032, AW974786, AW975154, AW975002, AW971403, AW975965, AW969673, AW975019, AW976024, AW979204, AW877209, AW979098, AW975031, AW975105, AW979002, AW979134, AL119457, AW975028, AW972845, AW969816, AW969680, AW974964, AW969791, AW975942, AW972377, AW974998, AW974658, AW973717, AW975876, AW975959, AW974801, AW979090, AW973213, AW972880, AW969839, AW975930, AW975966, AW975952, AW971732, AW971375, AW979238,

HTTSC95	33	997589	1 - 447	15 - 461	BE903286, BE738224, BG028767, BF795719, BF692544, BG165712, BG178489, AA776488, AW341060, AI696936, AW674401, AV755763, BF036130, BE894799, AA599100, AI444996, AV755340, BE617710, BE72009, AI133293, BF690592, BE748498, BG106158, BE900415, BG170173, BG111506, BF691436, BF667047, BF128835, BF527656, BF984042, BG106739, BF978871, BF312361, AV685933, BG166997, BE781635, BF698657, BF978019, BE277902, BE387231, BE892002, BF674900, BF244511, BF978866, BE297792, BE548639, BE730193, BF826005, BE540782, BF697609, BF796466, BG164429, AC005569, M23613, M28699, M26697, X16934, L15317, L15320, AC005192, AF164686, M81257, AC004941, AF164690, AL353580, M25062, U70984, AF164687, M33212, L15316, AC021037, AC021019, AC007068, AF133299, AF285089, AF285087, AC004994, AL023803, AF195956, AL035588, X54171, AC004538, L15318, L15319, AF164689, AC007349, J04943, AK000472, U41742, D45915, U41743, AF285088, X17200, L15315, J04944, M31004, J02590, U04946, U89321, U89311, M37041, M37036, AC010186, and U89309.
HTTIS02	34	999204	1 - 927	15 - 941	BF946981, AA077171, D17516, AR028892, AR028900, AR028894, AR028893, AR028902, AR028901, AR028895, AR028903, AC006466, AR028890, Z23279, AR028898, D14908, AR028889, L16506, AR028891, D82935, AR028897, Z23275, Z23274, Z23272, Z23282, AR028899, D14909, Z23273, D16465, L16680, AR028888, and D17290.
HTXQD18	35	1000599	1 - 565	15 - 579	AL046307, Z78386, AL572458, F27151, AA055750, AA333594, BE882817, AW778731, AW015540, BE222903, AU148865, AA099751, BG109260, U80751, AA716626, AW998255, BE267723, BE560019, AA040821, AA397648, AU130210, BE929525, BE538666, AB014542, AK001739, U90567, and AK027021.
HKGAW0 2	36	1003346	1 - 519	15 - 533	AU123654, BE622339, AA160849, BE872390, AW993641, AW840238, AA161281, AI755276, BE313918, AL525283, BF737162, BF038800, BE259923, AW068564, BF343512, AL492558, BF740307, T18598, BF083151, AA027021, AI372674, BF370280, BE168407, AW965680, BE707955, BG015537, AA143489, BE146202, BE868734, AK001199, AK024649, AB037858, and AX011688.
					AA935168, AW970942, AA398801, AW971975, AW972296, AW972786, AW975941, AW973101, AW972154, AW979160, AW979208, AW973190, AW975981, AW973821, AW969772, AW972817, AW969694, AW971326, AW975971, AW979147, AW973805, AW971953, AW974806, AW975166, AW971404, AW976024, AW975596, AW979204, AW972440, AW973812, AW973789, AW971403, AW979076, AW973254, AW971954, AW973762, AL119484, AW975434, AW973278, AW973252, AW970936, AL134524, AW975626, AW975161, AW975952, AW979212, AL119418, AW974658, AW971968, AW973819, AW979176, AW979211, AW974338, AW970969, BG059382, BG153750, AW975244, AW979037, AW979250, BE294931, AW973764, BE296530, AW972710, AW969930, AW979090, AW973825, AW974379, AL119483, AW973810, and AL135928.

HJRC82	30	993759	1 - 2034	15 - 2048	BE614945, AU123654, BE872390, AL525283, AW993641, AW965680, BF309693, BF308936, BF333907, BF737162, BE168022, AI492558, BE297137, AW068564, AI755276, BF818539, BG008965, AW840238, BF770908, AA160849, BE622339, AI372674, AA351821, BE168407, AA349466, BG015537, AI205125, BF769382, BF083151, BF511513, BE146202, BF873805, AW888417, AA161281, BF088733, T18598, BE707955, AA027021, BF927721, D31580, BF372972, BE003963, BF372969, BE313918, BF038800, R45887, BE259923, BF343512, BF740307, BE296215, BF927956, AB037858, AK024649, and AK001199.
HIABW64	31	994001	1 - 451	15 - 465	AW673312, AA310521, BG257922, AL522572, AU128744, AU137022, AA314039, BG176769, AV658758, AV650660, AA232595, AV691223, BE541064, AA223451, AA081255, C16722, AA188117, AK022951, AL353771, and AK000338.
HJBT01	32	997082	1 - 2339	15 - 2353	AL529942, AL537010, AL516075, AL515134, AL514852, AL516169, AL516168, AL518974, BF976161, AL517163, AL516116, AL517981, BE748906, BE748680, BE738582, BE739794, BE748858, BE748720, BE739931, AW027051, BE878041, BE748719, BE748689, BE748772, BE616122, BE748844, BE875343, BE616640, BE748801, AA603223, AI863055, BE748842, BG107746, AI818891, AA809143, AI887690, BG028229, BE739461, AW245991, AW080586, AA652496, AW275857, BE748564, BE616331, AV699966, AI718659, AI889997, BE748796, AW471400, AI609209, AI937155, BF732406, AI797841, BE793252, BE379655, AI815130, BE899125, AW166104, BE620337, AW192472, BG176573, BG106990, BE748575, BE275043, BE613215, AV688819, AW089551, BE734332, AI832414, BF966053, AI857822, BG113009, BE748833, BE748793, AW090035, BG232105, BE739818, BE798148, BE734941, AV716558, BE875560, BE748860, AI888796, BE267459, BE547819, BE250388, AW438831, BE383707, BG032786, BE796977, BG258965, AW514674, AW173432, BF828847, AV734973, BG109822, AI921611, AI889660, AA630772, AW612925, AI499137, BE738189, BE561714, AI814977, BE267271, AW167236, AI685210, BG109430, BE734983, AW512124, BG170680, AI340313, AL048498, BE744738, AW612903, AW886393, BF825320, AW512431, BE410410, BE897271, BE793623, AW571862, BG033255, AW071120, AL518973, BE540410, AW780430, BE793036, AV692453, AW440307, AW810385, BE622374, BE617121, BF978895, BE889388, BE221443, BG256591, BE900492, BE543722, BE748085, BG104559, AA729020, BE302279, BG106480, BF346781, BE748632, AW514653, BG254940, BE535876, BG033967, AW778862, BE867093, BF185080, AI619454, BE302284, BE350741, BE891310, BE734783, BG121347, BG258367, BE541541, AA595393, BF701162, AI559573, BE784652, BE786819, AW732803, BF061721, BF664133, BF240482, BE886973, BE792312, AV753308, BF026558, BE795433, BF694130, BE732901, AW471068, BF691650, BG257869, AW513494, AI571845, BE350331, BG032593, BF825795, AI568328, BE902403, AA482747, BE785524, BE877504, BE786682, BF026485, BE782431, AI811381, BE559901, BG256633, AI561091,

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HNGGK47	29	993602	1 - 686	15 - 700	<p>AA587109, AW813774, AW959611, AA149541, AI184943, AA283852, AA915990, AI754552, BE146143, BE146078, AA192458, AI276859, BF058388, AW968813, N25247, AA179831, R50526, BE906097, BE748870, AA489709, BF058508, BG001227, AA600248, AA149622, R50552, AI831786, H60862, BE748864, AA159263, AA912651, AV704542, AI360851, AI094704, AA151772, AA088441, AA130533, AI732442, AA130457, AA115727, AA132985, AA156548, AA100473, AA156552, AA133314, AA146791, AA115406, AA085900, AA506505, H30250, AA130431, AV705967, R50721, AA156558, AA593788, AA134890, AA160532, AA115791, AA056456, AI732411, AI144451, AW975853, AA837803, AI732210, AI732209, AA527289, BE145948, N43184, AA216639, AA233457, AA055823, AI535936, AI242047, BE146358, BG255562, AW817478, AA418333, AI264351, AI274932, Z99365, AA132266, AA155720, BE675378, H69025, AA483160, AA484178, AI685396, AI479610, AA152342, AA890486, BE146091, R78417, AA219599, AA223178, AI052113, AA100721, AA703512, AA213692, AA627379, R98355, AW593679, AA102795, AA227040, AW341960, H38219, T53321, W28465, AI762542, AA052890, AA242847, AA252138, AA134470, AA099427, AA129454, AA135403, AA102232, AA135390, AA689437, AA173762, AW129244, AW007800, AI017153, AA227186, AA548735, AA308744, AA223472, AA72499, AW854595, AA962219, AI636602, AI700160, H51291, AA055663, AA984688, AA100673, AA143593, AA528777, AA838754, AA434477, BE748279, AI949977, BF673870, AI280212, BG013985, AW969910, N66779, W25656, AW025824, AA424412, AI743039, AA135407, AW966820, BE741129, AI285813, AA405703, AA832265, AA157818, AA219559, BF816863, AA424458, AA936323,</p>

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HOFNI12	25	982193	1 - 471	15 - 485	X66139, and X66140.
HEGAZ61	26	983192	1 - 547	15 - 561	BF794968, BE396447, BE514415, BE513758, BE296708, BE878606, AW340333, AI806295, AW407893, AW268810, BF761940, BE268600, AA827664, AA829237, AA909185, BE243123, AA919008, AA604425, AI539428, AA011359, BF511512, AX027541, AB016590, AF241789, and AB016589.
HHERQ79	27	985266	1 - 749	15 - 763	BF972832, BF968664, BF337801, BF038410, BF338126, BF688743, BF688560, BF339410, BF726248, BE737528, BF338788, BE615509, BF528351, AI800218, AI346281, BG222425, BF036565, BF689301, AI983710, AW026429, AI766300, BF805547, BG249715, AI583539, AW264048, AI911743, BG060162, BG117232, AW387854, BE615228, AI924093, AI312053, AW470078, AI264211, AI955668, AI701482, BE677699, AW058431, BE045835, AW196781, BE221691, AI022218, BF983475, AI565141, BF689198, AI276048, BE736613, AI566956, AI689249, AI920904, BG056452, AW193090, AI623560, BE018907, BF570443, AW664907,
HRABV49	28	988947	1 - 2030	15 - 2044	

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HSIG160	24	956986	1 - 1316	15 - 1330	AW084660, AW074849, R52809, N51830, H11799, F06391, F05980, N45731, AI873913, BF946817, R11686, AV722459, N79408, H08526, AV722602, R51484, N79394, N54098, AV738340, AW964468, AW966389, AW949645, AW966330, D80045, AW965158, AW975618, AV702035, AW964532, AW949642, AV744012, AV742732, AV742050, AW949630, AV744690, AV704180, C14389, AW949643, AV706229, AV724520, AV720150, AV741220, C15076, AV705134, D59467, AV705869, AV742048, AV727978, D80164, D81026, AV718681, D59787, AV741191, AV706147, AV718692, AW973485, C14331, AV701004, D59859, AW366296, AV718489, AW973541, AW966053, AI535660, AV718530, AW978634, AV701643, C14429, AV719468, AV703542, D59889, AV741187, AV741198, AW949634, AV699479, AW959799, AW966041, AW965177, AW964488, AV699550, AW753053, D80024, D59275, AW966062, D80522, AV726812, AW966534, D80391, D81030, AW973330, AV720203, AW960465, AW966075, AW966065, AW966531, AW959469, AW949633, AV718770, AW966022, AV699652, AV706910, D58283, D59927, AW958992, AW959582, D51799, AW966013, D80196, D80022, D51423, D80166, AW978661, AV718800, D80195, D80253, AW975621, AV718440, AV720028, D59619, AW949654, D80210, AV699927, AW960553, D80240, AV719822, AV705453, AV720791, AW964541, AW966054, AW966029, AW966050, AV719188, AV699447, AW973307, D80043, D80227, AV719324, AV719783, D59502, AW959597, AV718633, AW959628, AW949641, AW975605, AW959570, AW973334, AV720211, AV718931, AV718844, AV720464, AV719557, AV720731, AV722801, AV723927, D59610, AW949656, AW959202, AW966043, AW956434, D80212, AI905856, AV720878, AW378532, AV744662, D80188, D50995, D51022, AW965163, AW966059, D80219, AW978648, AV702322, AW973474, D57483, AV704660, AW975613, D80366, AW966030, D80038, AW964756, D80269, AW958993, AV718938, AW960473, AW965185, AW965197, AW965196, AV718707, AW965184, AW973488, AW965175, AW973482, AW959136, AW959062, AW964477, AW962082, D50979, D80193, AW375405, AW973447, D80251, AW949657, AV720533, AW962245, AV706028, AV728436, AW949629, D80241, AW949653, AW949646, AV707151, AW949632, AW949658, AW949631, AW949618,

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HDQES11	21	950838	1 - 2069	15 - 2083	AL040398, AL043338, AI917735, BG163397, BE733691, AW303607, BG259714, AI819365, BG165205, BE888665, BE388695, BF694553, BE789604, AI243857, AW958665, AA496257, AW469326, AA278626, AA278625, AW469635, AA417787, AA465309, AA306656, AA831859, AA769522, AA312210, AA157110, AA579065, AA564476, AI476034, AI205635, AA740386, W23045, AA491643, AW995688, AA810400, F29732, AA417708, BE677853, AI567332, AW364598, BE148761, BE739252, BF830136, BE703932, BE831081, BG259221, BE703947, BE703993, BE831119, BE799580, AF221636, AF221635, AF221634, AF221637, and AK000290.
HOHCE96	22	955622	1 - 890	15 - 904	BE082811, BF368839, AA368624, BG003457, AA368519, AA853078, R30943, and AF206329.
HBKDR69	23	956894	1 - 1032	15 - 1046	BF574365, AI659815, BF002896, BF575970, BF575494, BF790003, BF790843, BF574388, BF574862, BF126301, BF436949, BF695348, BF671093, BF790369, BF573998, BF671272, AI458091, BF589170, AI675624, AI554082, AA211590, AI375478, AW016729, AA026659,

					AFI19909, AFI30110, AFI118094, A08916, A08913, AL359941, AK000323, AK025084, AF090934, AF158248, AF177401, AK026592, AK026480, AL117457, Y16645, AL353957, AF090886, AL050116, AFI16631, AK000718, AFI04032, A08910, AK024538, A08909, AK027204, AFI30105, AL359601, AFI30055, AR079032, AL133075, AL137526, AL137459, AK026865, AK000652, AK024524, AFI13694, AK027096, AFI19899, AFI19875, I89931, AL117460, AL050149, AL133016, AL137527, X65873, AF314091, AK027113, AFI16688, AL049938, AR087170, AK026959, AF079763, AL133560, AFI13019, AFI113677, AR070212, AX019230, AB052191, AL137550, AL122118, AFI30104, AB048953, AL133080, AFI119878, AL049382, AL122098, AF090900, AFI30077, AL389982, AF078844, E06743, AF017152, AL162062, AFI16682, AX019229, AK024992, AL359618, AK026784, AK027164, AFI13013, AF061795, AF090903, AFI51685, AK026532, AL122121, AL049430, AK026045, AFI83393, X84990, AF321617, AFI06862, AL137479, AR011880, AK000250, AF242189, AK025383, AF097996, E05822, I33392, AL137271, AK026744, AK026506, AK000083, AL162006, AK026927, AL133568, AK026528, AFI32676, AB047904, AL133640, AF061836, AFI13222, AR059958, AFI11849, U96683, AFI130087, I09499, AFI25948, AL389939, I00734, AL110225, AL133606, A93016, E03348, AFI17437, AL049452, AFI32205, A77033, A77035, AK026885, AL137529, Y09972, AK025708, AFI113676, E00617, E00717, E00778, X82434, AL122049, E04233, AL122050, AK000137, AL133557, AFI116602, AL050393, AK026534, AR038854, AFI116644, AL137557, AB051158, Y11254, AL110196, AK026583, AFI11851, Z82022, AL110221, AC002467, AB047887, AFI13699, E02349, AK026542, AK026741, AB048954, AK025092, AFI30082, AF207829, AFI19894, AK027081, AFI118064, AJ000937, AB047615, AFI19865, AL133081, AFI77336, AL359583, AL117583, AK027146, AK025524, AK025772, AL096744, AL050146, AL050108, Y07905, AK026855, AF008439, X72889, AFI16691, AB052200, AK026504, AK026642, AK026600, AFI146568, S79832, AF022363, AL122123, I03321, AFI18070, AB050510, S78214, and AFI38861.
HWLHQ59	20	949724	1 - 1281	15 - 1295	AW842606, AW842562, AL134524, AW877209, AW975037, AW979127, AW979134, AW979098, AL119324, AW972292, AW976024, AW975002, AW861944, AW979204, AW975032, AW858525, AW975965, AW858526, AW969791, AW975105, AW971403, AW975954, AW971975, AW975942, AW975031, AW975019, AW975649, AW975028, AW971404, AW974786, AW979238, AW972849, AW975876, AW858522, AW975154, AW979090, AW979176, AW975230, AW976023, AW969673, AW971968, AW969885, AW975971, AW972377, AW975015, AW979106, AW974975, AW970945, AW969680, AW975981, AW973219, AW979294, AW969816, AW975027, AW979212, AW974964, AW973750, AW975632, AW979054, AW974823, AW970942, AW975020, AW972680, AW858455, AW975025, AW975959, AW974785, AW975244, AW975952, AW974998,

AA978174, AI205675, AI183657, AW954031, AW958647, AV709869, AW963492, AW961037, AW949453, AV693527, AV651436, AV702932, AV707296, AV728806, AV727512, AW949931, AV727990, AV702117, AV725055, AV654896, AW952439, AV685966, AW950438, AV727839, AW949433, AV645906, AV707727, AI525669, AV705341, AV707024, AV708622, AV701620, AV704182, AV683313, AW960594, AV708834, AV726605, AV704934, AV690007, AV650703, AV703472, AV708075, AV686346, AI525653, AV683272, AW945168, AV704696, AV707889, AV705323, AV729189, AV728670, AV726784, AV707767, AV702833, AV661704, AV702147, AW961388, AW963481, AV703695, AV702994, AV655425, AW952457, AV708723, AV708653, AV698087, AV728592, AW953787, AW953965, AV722222, AV753074, AV703042, AV708980, AV729462, AV721957, AV709604, AV652027, AV705384, AV706185, AV705632, AV708004, AW966420, AV704955, AV701560, AV705693, AW951263, AV689111, AV728157, AW959910, AV647118, AV682466, AV726951, AV687535, AV726229, AW960326, AW955653, BG109125, AV703970, AW965144, AV656903, AV706721, AV655280, AV706987, AV727980, AV729451, AV702430, AV655473, AV660515, AV708025, AV707274, AV727776, AW950035, AV701580, AW951273, AW953790, AW951618, AV692691, AV706775, AV729259, AV704824, AV687035, AW965502, AV704269, AV708992, AV726738, AV660662, AV705280, AW953792, AV703372, AV693985, AV704345, AV696106, AV728518, AV698043, AV650024, AV703168, AV682502, AI811344, AI677796, AW959837, AV708704, AV725580, AV708438, AV729439, AV654624, AV652156, AV703263, AV707753, AW961463, AV703222, AV651983, AW953967, BF814447, BF032768, AW965840, BF338002, AV659536, AV709517, AV652443, AV702804, AV659322, AI513999, AW955606, AW949732, BF814450, AV705938, AV708893, AV705348, AV703362, AV648430, AV726103, BF344031, AI541056, AI499285, AV696257, BF968653, AW960720, AI802542, AV697196, AI815855, AV652001, BG179993, AV660592, BF970449, AV709390, AV703495, AI921176, AI564719, BF822127, AV656478, AV708381, AV660728, BF793309, AI040243, AI358701, AV725920, BF882334, AV706744, AV706915, AV702021, AV715560, AV704350, AI541048, AV726624, AV655096, BG121959, AV701707, AV652617, AV648263, BF037484, BE910373, AI611743, AI619502, BG249582, AI816010, AW161579, AV709314, AV658835, AV684417, AI633125, AI039276, BE880182, BE047606, AV727799, AV729263, AV728733, AV703169, BF792961, BF526020, AI284484, AI884469, BF970768, AV705223, BF970652, BF037097, AI921248, BF814420, AV684604, AV727029, AV647659, BG029033, AI433157, AI133004, Y08617, I48978, S68736, AF218031, X70685, I48979, AB048975, AI162083, AK025339, AF090901, AK025414, A65341, I89947, AK026630, AK025967, AK024588, AB041801, AI116646, AI1353940.
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					AC005658, AL117258, AP001705, AC005095, AL139383, AC020728, AC006582, AC007773, AL136162, AC007450, AP000108, AP000040, AF111169, AL161657, D84394, AC004707, AC006387, AC012064, AC007221, AC004838, AC005102, AC005036, AC019210, AC006088, AP001746, AC009946, AC008072, AC022392, AL161422, AC002331, AC011456, AP000240, AC020552, AC004964, AL138812, AP001715, AC073316, AL023693, AL139350, AP001684, AL049541, AC015555, AC005534, AL035608, Z92546, AP000353, AC013434, AC018731, AC005295, AL133383, AP000953, AC004910, Z98257, AC005659, AP001694, AL163280, AC002992, AL359554, AC006241, AL121657, AL034429, AC004510, AC005548, AP001207, AC007376, AC009501, AL162293, AC005993, AL031681, AC004825, AC004976, AL137786, AC004409, AC004896, AC022206, AL359218, AL078646, AC007251, AL162584, AC004859, AL157398, AL096776, AL096816, AC019187, AL133346, AL133391, AB020863, AL139352, AL356276, AC004525, AL137022, AC012067, AC002464, AC009429, AC006167, AC009086, AC002390, AL031658, AL021806, AC005047, AP001710, AC002538, AC010150, AL353748, AB020868, AC004887, AC011604, AP000284, AC004542, AL031291, AP000283, AC005261, AC005477, AL442166, AL031772, AL121578, AC011311, AL163284, AL031676, AL445466, AC005027, AL035420, AC004817, AF096876, AC010282, AF029308, AC073323, Z93015, AC022073, AL160234, AC007240, AC015973, AC008083, AP002898, AL121755, AL137800, AF311103, AL035408, Z93942, AL137145, AL031663, AC006536, AC008764, AL133289, AC008180, AL136137, AC007739, AL157361, AC010463, AP000511, AL121675, AC009484, AC007402, AC005547, AC008536, AC005838, AB038162, AL109855, Z82245, AP001208, AP001707, AR036572, U91328, AC005616, AC008750, and AC008750.
HTPAU91	13	846654	1 - 2463	15 - 2477	AA295383, and AL355178.
H7TBE17	14	858079	1 - 692	15 - 706	BF900226, AR075074, AR075073, AL355388, and AL355388.
H7TRA52	15	858086	1 - 691	15 - 705	BF900226, AR075074, AR075073, AL355388, and AL355388.
HSSIC65	16	872582	1 - 1760	15 - 1774	BE889609, BF183110, BG248693, AA452195, AI821464, AI621234, AI821685, AI745288, AI655244, AA436362, BF432211, AW770404, BF196926, AI820935, AA427404, H08678, AI432244, AA428418, H08677, AA410413, H61387, AA410412, H14737, BF351660, BG170733, H62405, AA410253, AI968569, AW002476, AW166810, BE350305, BF530203, H09849, AW748526, AX047642, AC058790, AC007663, AC006549, AB045987, and AC006549.
HKLAA30	17	875001	1 - 398	15 - 412	AV655586, AL527281, BE313499, AV690347, BE314871, BE259942, BF314316, BE260383, AL527320, BF312192, AA089855, H30455, AA954657, and AA455419.
HPJCP59	18	943763	1 - 1069	15 - 1083	AW297778, BE831642, AA704347, AI693619, AF269255, AL133353, and AF288221.
HTEB072	19	947922	1 - 791	15 - 805	AI073398, AA906424, AA724715, AW665390, AA969255, AI149619, BE866717,

					AL024498, AP000514, Z98200, AC004971, AC005280, AC018663, AL050307, AL049872, AC008556, AC022432, AC011500, AL445263, AP01628, AL021808, AC005740, AC006057, AC008940, AC002091, AC007664, U78027, AC002306, AL031846, AL136228, AL049709, AL138725, AC006581, AL356299, AC005098, AC004847, AL355610, AL021453, AC008738, AL365315, AC011490, AL022323, AC016683, AC009362, AL139188, Z83844, AL117186, AF129756, AC007057, AB039887, L78833, AC006483, AL096764, AL117377, AC006538, AL034379, AL035422, AC025593, AC006511, AL138756, AL035089, AC005365, AL158040, AL135744, AC005527, U93163, U91327, AC008952, AC005920, AC006449, AJ271735, AC006162, AL117334, AC008773, AL035696, AL354836, AC008626, AC007637, AC002137, AC005003, AC002045, AJ011930, AJ003147, AC008155, AL353807, AC007050, AL031257, AC006013, AC008018, Z97630, AC010422, AC005257, AC007889, AB011399, Z98752, AL133174, AF001549, AC010458, AC005103, AP000248, AC005589, AL353748, AL121655, AF047825, AC009756, AC006512, AL163300, AF241734, AC005618, U85195, AL356532, AC010789, Z95116, AC007389, and Z83822.
HDFMQ69	12	812744	1 - 3355	15 - 3369	BF663289, AI880327, AW513296, BF893706, AL766223, BE043031, BG032943, AI811827, AW088758, AW812622, AI948443, AI821377, N20948, AI690026, AL042906, AA680243, BG260565, AA321342, AI918403, AU119532, AI223036, BF816106, AV760760, AW504485, AI732120, AU140403, AA433949, AI978654, BF527620, N35317, AL042905, AV733710, AA613164, AV762220, AV760723, AA621333, AL043745, T64484, BF872337, AV761207, BE883679, AI734144, AA524604, AW172908, BF982222, AI744949, AV761013, BE740358, AW962035, AW962925, BE675240, AI681189, AV704568, AW963463, AA487142, AA410788, BF816974, AI474187, AI627917, AV763538, AV757289, F03312, AA535216, AL079734, AI821044, BF936016, BF826830, AU140392, AW973406, AL138254, BF736198, BE062476, AI609972, AV762783, AW409807, AW820439, AL042373, BE327876, AV759632, BG029528, AW816516, AW969667, AC008750, AC011452, AC018755, AC020914, AC005011, AP001053, AC010358, AC006480, AC006337, AP001752, AC007263, AL049872, AC011490, AC005874, AF134471, AC010510, AC026704, AC004965, AL157414, AC008784, AC026122, AL117338, AC007225, AC005228, AC004702, AC024952, AC007536, AC005799, AC007546, AC004791, AL109806, AL031767, AC005482, AC007685, AC004980, AF001549, AC005953, AC006509, AC004067, AL158172, AL136418, AC083866, AC023051, AC010485, AC016594, U82670, AC005207, AC005522, AC002375, AC010473, Z97055, U52112, AC011093, AC010206, AC009475, AC007510, AF001708, AC004921, AL035414, AC004605, AL138755, AB023050, AP000509, AC007276, AC004185, AL109918, AC020946, AC083864, AC004797, AL139824, AC025594, AF172277, U91318, AC009060, AF207955, AC006059, AL132774, AC003037, AP001623, AL162615, AL355101, AC008651, AL133355, AL136969,

TABLE 3

Clone ID NO: Z	SEQ ID NO: X	Contig ID:	EST Disclaimer Range of a Range of b		Accession #'s
			1 - 2006	15 - 2020	
HEMEC34	11	684296			AW410844, BG180320, AW265468, AI890283, AI064968, AI826857, AI702049, AV763460, AI753131, AV758849, AI732720, AI444575, BG059139, AL121039, AI754257, AW504667, AW970856, AW021674, BG250794, AW148821, AA569220, AA640305, AI572680, AF126947, AI567676, BE244308, AA935827, BE080768, AI744199, AW243817, AW023390, BF973510, BF949151, AI921744, AV734311, AW873417, AI753904, AA595661, AW069110, AA330549, AI568376, AA489390, BG109302, AA610644, BG112045, AI570067, AW192930, AW020682, AI828721, AI751698, AI434103, AI609992, AI312267, BG004739, AL138262, BG059924, AV741384, AI445699, AV731938, AW192419, BE045167, AA55232, AW675677, BF679568, AW968167, AI308529, AV756809, R61887, AI734119, BG169186, BF736912, AV712092, AA313025, AI345166, AA834891, BF849260, AI252611, AA055647, AI167715, AI792586, AT92498, AV759354, AA809546, AI884404, AI701898, BE139452, AF039390, AL133412, AL022315, AC005565, AL118506, AC005519, AL121712, AL096840, AF207550, AC005071, AC018641, AC005080, AF196779, AL049760, AL035460, AC011475, AF134726, AC008044, AP001711, AL139353, AF081786, AL109952, AP000511, AL136304, AB023048, AC016026, AC009470, AL035587, AC005821, AD000092, AL121872, AP000030, AC006315, AF075069, AC005629, AC026776, AC004263, AP000503, AP000211, AP000133, AF196971, AC000159, Z73979, AC005971, AL136418, D00591, AL135787, AP001712, AC000115, Z97632, AL139389, AC006050, AC010627, AC010271, L44140, AC006139, AL031657, AC007684, AC003982, AL121886, AC007782, Z93020, AL139343, AC005937, AC007956, AL080317, Z99714, AC008649, AC000070, AL022726, AC000082, AC004673, AL137230, AP000694, AL118499, AC004876, AC006453, AC009086, AC012309, AL035086, AC008752, AL121653, AL139099, AP001725, AC002991, AC004019, AC002310, AL133163, AC011465, AL031670, AL035416, AC006064, AL360157, AL132866, AC020904, AC000353, AC002543, AL121601, AL122001, AC005484, AC008277, AP001629, AL133387, AC005932, AF051976, AC027319, AC027612, AL117381, AC007845, AC007731, AL034549, AC005500, AC005180, AL031311, AC004821, AC005325, AC018645, AL031587, AC021752, AC016554, AL354948, AF225899, AC005412, AC016025, AL133545, AL050349, AC004999, AC020934, AI295844, AC004765, AC009032, U80017,

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contig sequence (SEQ ID NO:X) listed in the fourth column of Table 1, preferably excluded are one or more polynucleotides comprising a nucleotide sequence described by the general formula of a-b, where a is any integer between 1 and the final nucleotide minus 15 of SEQ ID NO:X, b is an integer of 15 to the final nucleotide of SEQ ID NO:X, where both a and b correspond to the positions of nucleotide residues shown in SEQ ID NO:X, and where b is greater than or equal to a + 14. More specifically, preferably excluded are one or more polynucleotides comprising a nucleotide sequence described by the general formula of a-b, where a and b are integers as defined in columns 4 and 5, respectively, of Table 3. In specific embodiments, the polynucleotides of the invention do not consist of at least one, two, three, four, five, ten, or more of the specific polynucleotide sequences referenced by the Genbank Accession No. as disclosed in column 6 of Table 3 (including for example, published sequence in connection with a particular BAC clone). In further embodiments, preferably excluded from the invention are the specific polynucleotide sequence(s) contained in the clones corresponding to at least one, two, three, four, five, ten, or more of the available material having the accession numbers identified in the sixth column of this Table (including for example, the actual sequence contained in an identified BAC clone). In no way is this listing meant to encompass all of the sequences which may be excluded by the general formula, it is just a representative example. All references available through these accessions are hereby incorporated by reference in their entirety.

[0088] The polypeptides may be in the form of the secreted protein, including the mature form, or may be a part of a larger protein, such as a fusion protein (see below). It is often advantageous to include an additional amino acid sequence which contains secretory or leader sequences, pro-sequences, sequences which aid in purification, such as multiple histidine residues, or an additional sequence for stability during recombinant production.

[0089] The polypeptides of the present invention are preferably provided in an isolated form, and preferably are substantially purified. A recombinantly produced version of a polypeptide, including the secreted polypeptide, can be substantially purified using techniques described herein or otherwise known in the art, such as, for example, by the one-step method described in Smith and Johnson, Gene 67:31-40 (1988). Polypeptides of the invention also can be purified from natural, synthetic or recombinant sources using techniques described herein or otherwise known in the art, such as, for example, antibodies of the invention raised against the polypeptides of the present invention in methods which are well known in the art.

[0090] The present invention provides a polynucleotide comprising, or alternatively consisting of, the nucleic acid sequence of SEQ ID NO:X, and/or the cDNA sequence contained in Clone ID NO:Z. The present invention also provides a polypeptide comprising, or alternatively, consisting of, the polypeptide sequence of SEQ ID NO:Y, a polypeptide encoded by SEQ ID NO:X or a complement thereof, and/or a polypeptide encoded by the cDNA contained in Clone ID NO:Z. Polynucleotides encoding a polypeptide comprising, or alternatively consisting of the polypeptide sequence of SEQ ID NO:Y, a polypeptide encoded by SEQ ID NO:X, and/or a polypeptide encoded by the cDNA contained in Clone ID NO:Z are also encompassed by the invention. The present invention further encompasses a polynucleotide comprising, or alternatively consisting of, the complement of the nucleic acid sequence of SEQ ID NO:X, a nucleic acid sequence encoding a polypeptide encoded by the complement of the nucleic acid sequence of SEQ ID NO:X, and/or the cDNA contained in Clone ID NO:Z.

[0091] Many polynucleotide sequences, such as EST sequences, are publicly available and accessible through sequence databases and may have been publicly available prior to conception of the present invention. Preferably, such related polynucleotides are specifically excluded from the scope of the present invention. Accordingly, for each

[0084] Vectors pSport1, pCMVSPORT 1.0, pCMVSPORT 2.0 and pCMVSPORT 3.0, were obtained from Life Technologies, Inc., P. O. Box 6009, Gaithersburg, MD 20897. All Sport vectors contain an ampicillin resistance gene and may be transformed into *E. coli* strain DH10B, also available from Life Technologies. See, for instance, Gruber, C. E., et al., *Focus* 15:59- (1993). Vector lacmid BA (Bento Soares, Columbia University, New York, NY) contains an ampicillin resistance gene and can be transformed into *E. coli* strain XL-1 Blue. Vector pCR[®]2.1, which is available from Invitrogen, 1600 Faraday Avenue, Carlsbad, CA 92008, contains an ampicillin resistance gene and may be transformed into *E. coli* strain DH10B, available from Life Technologies. See, for instance, Clark, J. M., *Nuc. Acids Res.* 16:9677-9686 (1988) and Mead, D. et al., *Bio/Technology* 9: (1991).

[0085] The present invention also relates to the genes corresponding to SEQ ID NO:X, SEQ ID NO:Y, and/or the deposited clone (Clone ID NO:Z). The corresponding gene can be isolated in accordance with known methods using the sequence information disclosed herein. Such methods include preparing probes or primers from the disclosed sequence and identifying or amplifying the corresponding gene from appropriate sources of genomic material.

[0086] Also provided in the present invention are allelic variants, orthologs, and/or species homologs. Procedures known in the art can be used to obtain full-length genes, allelic variants, splice variants, full-length coding portions, orthologs, and/or species homologs of genes corresponding to SEQ ID NO:X or the complement thereof, polypeptides encoded by genes corresponding to SEQ ID NO:X or the complement thereof, and/or the cDNA contained in Clone ID NO:Z, using information from the sequences disclosed herein or the clones deposited with the ATCC. For example, allelic variants and/or species homologs may be isolated and identified by making suitable probes or primers from the sequences provided herein and screening a suitable nucleic acid source for allelic variants and/or the desired homologue.

[0087] The polypeptides of the invention can be prepared in any suitable manner. Such polypeptides include isolated naturally occurring polypeptides, recombinantly produced polypeptides, synthetically produced polypeptides, or polypeptides produced by a combination of these methods. Means for preparing such polypeptides are well understood in the art.

preparation can then be used as a template for first strand cDNA synthesis using a gene specific oligonucleotide. The first strand synthesis reaction can then be used as a template for PCR amplification of the desired 5' end using a primer specific to the ligated RNA oligonucleotide and a primer specific to the known sequence of the gene of interest. The resultant product is then sequenced and analyzed to confirm that the 5' end sequence belongs to the relevant gene.

[0082] The present invention also relates to vectors or plasmids which include such DNA sequences, as well as the use of the DNA sequences. The material deposited with the ATCC (deposited with the ATCC on September 12, 2000 and given ATCC Designation Numbers PTA-2448 and PTA-2449) is a mixture of cDNA clones derived from a variety of human tissues and cloned in either a plasmid vector or a phage vector. These deposits are referred to as "the deposits" herein. The deposited material includes cDNA clones corresponding to SEQ ID NO:X described, for example, in Table 1 (Clone ID NO:Z). A clone which is isolatable from the ATCC Deposits by use of a sequence listed as SEQ ID NO:X, may include the entire coding region of a human gene or in other cases such clone may include a substantial portion of the coding region of a human gene. Furthermore, although the sequence listing may in some instances list only a portion of the DNA sequence in a clone included in the ATCC Deposits, it is well within the ability of one skilled in the art to sequence the DNA included in a clone contained in the ATCC Deposits by use of a sequence (or portion thereof) described in, for example Tables 1 or 2 by procedures hereinafter further described, and others apparent to those skilled in the art.

[0083] Vectors Lambda Zap (U.S. Patent Nos. 5,128,256 and 5,286,636), Uni-Zap XR (U.S. Patent Nos. 5,128, 256 and 5,286,636), Zap Express (U.S. Patent Nos. 5,128,256 and 5,286,636), pBluescript (pBS) (Short, J. M. et al., *Nucleic Acids Res.* 16:7583-7600 (1988); Alting-Mees, M. A. and Short, J. M., *Nucleic Acids Res.* 17:9494 (1989)) and pBK (Alting-Mees, M. A. et al., *Strategies* 5:58-61 (1992)) are commercially available from Stratagene Cloning Systems, Inc., 11011 N. Torrey Pines Road, La Jolla, CA, 92037. pBS contains an ampicillin resistance gene and pBK contains a neomycin resistance gene. Phagemid pBS may be excised from the Lambda Zap and Uni-Zap XR vectors, and phagemid pBK may be excised from the Zap Express vector. Both phagemids may be transformed into *E. coli* strain XL-1 Blue, also available from Stratagene.

the first-strand cDNA. This obviates the necessity for the dA-tailing reaction which results in a polyT stretch that is difficult to sequence past.

[0080] An alternative to generating 5' or 3' cDNA from RNA is to use cDNA library double-stranded DNA. An asymmetric PCR-amplified antisense cDNA strand is synthesized with an antisense cDNA-specific primer and a plasmid-anchored primer. These primers are removed and a symmetric PCR reaction is performed with a nested cDNA-specific antisense primer and the plasmid-anchored primer.

RNA Ligase Protocol For Generating The 5' or 3' End Sequences To Obtain Full Length Genes

[0081] Once a gene of interest is identified, several methods are available for the identification of the 5' or 3' portions of the gene which may not be present in the original cDNA plasmid. These methods include, but are not limited to, filter probing, clone enrichment using specific probes and protocols similar and identical to 5' and 3' RACE. While the full length gene may be present in the library and can be identified by probing, a useful method for generating the 5' or 3' end is to use the existing sequence information from the original cDNA to generate the missing information. A method similar to 5' RACE is available for generating the missing 5' end of a desired full-length gene. (This method was published by Fromont-Racine et al., Nucleic Acids Res., 21(7):1683-1684 (1993)). Briefly, a specific RNA oligonucleotide is ligated to the 5' ends of a population of RNA presumably containing full-length gene RNA transcript and a primer set containing a primer specific to the ligated RNA oligonucleotide and a primer specific to a known sequence of the gene of interest, is used to PCR amplify the 5' portion of the desired full length gene which may then be sequenced and used to generate the full length gene. This method starts with total RNA isolated from the desired source, poly A RNA may be used but is not a prerequisite for this procedure. The RNA preparation may then be treated with phosphatase if necessary to eliminate 5' phosphate groups on degraded or damaged RNA which may interfere with the later RNA ligase step. The phosphatase if used is then inactivated and the RNA is treated with tobacco acid pyrophosphatase in order to remove the cap structure present at the 5' ends of messenger RNAs. This reaction leaves a 5' phosphate group at the 5' end of the cap cleaved RNA which can then be ligated to an RNA oligonucleotide using T4 RNA ligase. This modified RNA

[0078] Partial cDNA clones can be made full-length by utilizing the rapid amplification of cDNA ends (RACE) procedure described in Frohman, M.A., et al., Proc. Nat'l. Acad. Sci. USA, 85:8998-9002 (1988). A cDNA clone missing either the 5' or 3' end can be reconstructed to include the absent base pairs extending to the translational start or stop codon, respectively. In some cases, cDNAs are missing the start codon of translation, therefor. The following briefly describes a modification of this original 5' RACE procedure. Poly A+ or total RNA is reverse transcribed with Superscript II (Gibco/BRL) and an antisense or complementary primer specific to the cDNA sequence. The primer is removed from the reaction with a Microcon Concentrator (Amicon). The first-strand cDNA is then tailed with dATP and terminal deoxynucleotide transferase (Gibco/BRL). Thus, an anchor sequence is produced which is needed for PCR amplification. The second strand is synthesized from the dA-tail in PCR buffer, Taq DNA polymerase (Perkin-Elmer Cetus), an oligo-dT primer containing three adjacent restriction sites (XhoI, SalI and ClaI) at the 5' end and a primer containing just these restriction sites. This double-stranded cDNA is PCR amplified for 40 cycles with the same primers as well as a nested cDNA-specific antisense primer. The PCR products are size-separated on an ethidium bromide-agarose gel and the region of gel containing cDNA products the predicted size of missing protein-coding DNA is removed. cDNA is purified from the agarose with the Magic PCR Prep kit (Promega), restriction digested with XhoI or SalI, and ligated to a plasmid such as pBluescript SKII (Stratagene) at XhoI and EcoRV sites. This DNA is transformed into bacteria and the plasmid clones sequenced to identify the correct protein-coding inserts. Correct 5' ends are confirmed by comparing this sequence with the putatively identified homologue and overlap with the partial cDNA clone. Similar methods known in the art and/or commercial kits are used to amplify and recover 3' ends.

[0079] Several quality-controlled kits are commercially available for purchase. Similar reagents and methods to those above are supplied in kit form from Gibco/BRL for both 5' and 3' RACE for recovery of full length genes. A second kit is available from Clontech which is a modification of a related technique, SLIC (single-stranded ligation to single-stranded cDNA), developed by Dumas et al., Nucleic Acids Res., 19:5227-32 (1991). The major differences in procedure are that the RNA is alkaline hydrolyzed after reverse transcription and RNA ligase is used to join a restriction site-containing anchor primer to

useful for designing nucleic acid hybridization probes that will detect nucleic acid sequences contained in SEQ ID NO:X or the cDNA contained in Clone ID NO:Z. These probes will also hybridize to nucleic acid molecules in biological samples, thereby enabling immediate applications in chromosome mapping, linkage analysis, tissue identification and/or typing, and a variety of forensic and diagnostic methods of the invention. Similarly, polypeptides identified from SEQ ID NO:Y may be used to generate antibodies which bind specifically to these polypeptides, or fragments thereof, and/or to the polypeptides encoded by the cDNA clones identified in, for example, Table 1.

[0075] Nevertheless, DNA sequences generated by sequencing reactions can contain sequencing errors. The errors exist as misidentified nucleotides, or as insertions or deletions of nucleotides in the generated DNA sequence. The erroneously inserted or deleted nucleotides cause frame shifts in the reading frames of the predicted amino acid sequence. In these cases, the predicted amino acid sequence diverges from the actual amino acid sequence, even though the generated DNA sequence may be greater than 99.9% identical to the actual DNA sequence (for example, one base insertion or deletion in an open reading frame of over 1000 bases).

[0076] Accordingly, for those applications requiring precision in the nucleotide sequence or the amino acid sequence, the present invention provides not only the generated nucleotide sequence identified as SEQ ID NO:X, and a predicted translated amino acid sequence identified as SEQ ID NO:Y, but also a sample of plasmid DNA containing cDNA Clone ID NO:Z (deposited with the ATCC on September 12, 2000 and given ATCC Designation Numbers PTA-2448 and PTA-2449). The nucleotide sequence of each deposited clone can readily be determined by sequencing the deposited clone in accordance with known methods. Further, techniques known in the art can be used to verify the nucleotide sequences of SEQ ID NO:X.

[0077] The predicted amino acid sequence can then be verified from such deposits. Moreover, the amino acid sequence of the protein encoded by a particular clone can also be directly determined by peptide sequencing or by expressing the protein in a suitable host cell containing the deposited human cDNA, collecting the protein, and determining its sequence.

RACE Protocol For Recovery of Full-Length Genes

of Query amino acids in the HSP and multiplying by 100. The polynucleotides of SEQ ID NO:X which encode the polypeptide sequence that generates an HSP are delineated by columns 8 and 9 of Table 2.

[0072] The PFAM database, PFAM version 2.1, (Sonnhammer et al., Nucl. Acids Res., 26:320-322, 1998)) consists of a series of multiple sequence alignments; one alignment for each protein family. Each multiple sequence alignment is converted into a probability model called a Hidden Markov Model, or HMM, that represents the position-specific variation among the sequences that make up the multiple sequence alignment (see, e.g., Durbin et al., *Biological sequence analysis: probabilistic models of proteins and nucleic acids*, Cambridge University Press, 1998 for the theory of HMMs). The program HMMER version 1.8 (Sean Eddy, Washington University in Saint Louis) was used to compare the predicted protein sequence for each Query sequence (SEQ ID NO:Y in Table 1) to each of the HMMs derived from PFAM version 2.1. A HMM derived from PFAM version 2.1 was said to be a significant match to a polypeptide of the invention if the score returned by HMMER 1.8 was greater than 0.8 times the HMMER 1.8 score obtained with the most distantly related known member of that protein family. The description of the PFAM family which shares a significant match with a polypeptide of the invention is listed in column 5 of Table 2, and the database accession number of the PFAM hit is provided in column 6. Column 7 provides the score returned by HMMER version 1.8 for the alignment. Columns 8 and 9 delineate the polynucleotides of SEQ ID NO:X which encode the polypeptide sequence which show a significant match to a PFAM protein family.

[0073] As mentioned, columns 8 and 9 in Table 2, "NT From" and "NT To", delineate the polynucleotides of "SEQ ID NO:X" that encode a polypeptide having a significant match to the PFAM/NR database as disclosed in the fifth column. In one embodiment, the invention provides a protein comprising, or alternatively consisting of, a polypeptide encoded by the polynucleotides of SEQ ID NO:X delineated in columns 8 and 9 of Table 2. Also provided are polynucleotides encoding such proteins, and the complementary strand thereto.

[0074] The nucleotide sequence SEQ ID NO:X and the translated SEQ ID NO:Y are sufficiently accurate and otherwise suitable for a variety of uses well known in the art and described further below. For instance, the nucleotide sequences of SEQ ID NO:X are

[0070] Table 2 further characterizes certain encoded polypeptides of the invention, by providing the results of comparisons to protein and protein family databases. The first column provides a unique clone identifier, "Clone ID NO:", corresponding to a cDNA clone disclosed in Table 1. The second column provides the unique contig identifier, "Contig ID:" which allows correlation with the information in Table 1. The third column provides the sequence identifier, "SEQ ID NO:", for the contig polynucleotide sequences. The fourth column provides the analysis method by which the homology/identity disclosed in the Table was determined. The fifth column provides a description of the PFAM/NR hit identified by each analysis. Column six provides the accession number of the PFAM/NR hit disclosed in the fifth column. Column seven, score/percent identity, provides a quality score or the percent identity, of the hit disclosed in column five. Comparisons were made between polypeptides encoded by polynucleotides of the invention and a non-redundant protein database (herein referred to as "NR"), or a database of protein families (herein referred to as "PFAM"), as described below.

[0071] The NR database, which comprises the NBRF PIR database, the NCBI GenPept database, and the SIB SwissProt and TrEMBL databases, was made non-redundant using the computer program nrdb2 (Warren Gish, Washington University in Saint Louis). Each of the polynucleotides shown in Table 1, column 3 (e.g., SEQ ID NO:X or the 'Query' sequence) was used to search against the NR database. The computer program BLASTX was used to compare a 6-frame translation of the Query sequence to the NR database (for information about the BLASTX algorithm please see Altshul et al., J. Mol. Biol. 215:403-410 (1990); and Gish and States, Nat. Genet. 3:266-272 (1993). A description of the sequence that is most similar to the Query sequence (the highest scoring 'Subject') is shown in column five of Table 2 and the database accession number for that sequence is provided in column six. The highest scoring 'Subject' is reported in Table 2 if (a) the estimated probability that the match occurred by chance alone is less than $1.0e-07$, and (b) the match was not to a known repetitive element. BLASTX returns alignments of short polypeptide segments of the Query and Subject sequences which share a high degree of similarity; these segments are known as High-Scoring Segment Pairs or HSPs. Table 2 reports the degree of similarity between the Query and the Subject for each HSP as a percent identity in Column 7. The percent identity is determined by dividing the number of exact matches between the two aligned sequences in the HSP, dividing by the number

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9						protein product [Homo sapiens]			37%	1	354
									30%	7	384

HNFO84	HNFO84R	177	blastx.2	precursor [Homo sapiens] unnamed protein product [unidentified]	emb CAB69577.1	48%	3	308
HNTRS86	HNTRS86R	178	blastx.2	platelet glycoprotein Ib alpha chain precursor [Homo sapiens]	gb AAAS2595.1	43% 58%	1 242	180 292
HOEAC11	HOEAC11R	179	blastx.2	(AF022813) tetraspan [Homo sapiens]	gb AAC51864.1	79%	145	261
HOFNH58	HOFNH58R	180	blastx.2	lysosomal membrane glycoprotein-2 [Homo sapiens]	gb AAA60383.1	62% 76%	232 180	558 218
HOHAL20	HOHAL20R	181	blastx.2	latent transforming growth factor beta-binding protein 3'1	pir A57293 A57293	61% 46% 31% 30% 35% 33% 56%	113 258 113 113 113 164 147	559 569 538 511 238 253 194
HS2AO61	HS2AO61R	182	blastx.2	EGF repeat transmembrane protein [Mus musculus]	gb AAB01338.1	77% 36% 39% 33%	39 42 60 75	335 326 266 326
HSIFN53	HSIFN53R	183	blastx.2	aminopeptidase N precursor (EC 3.4.11.2) [Homo sapiens]	gb AAA51719.1	57%	2	208
HWBEA69	HWBEA69R	184	blastx.2	HLA CLASS II HISTOCOMPATIBILITY ANTIGEN, GAMMA CHAIN 1	sp P04233 HG2A_ HUMAN	59% 100% 41%	259 210 380	351 251 430
HWCAA2	HWCAA29R	185	blastx.2	(AK001199) unnamed	dbj BAA91549.1	55%	1	354

HEMEP27	HEMEP27R	167	blastx.2	proline-rich protein MP2 - mouse (fragment)	pir A24264 A24264	41% 42% 41% 40% 41% 38% 41% 41% 36% 35% 39% 42%	6 6 6 6 6 6 6 4 6 6 4 5	305 281 281 305 281 323 260 303 281 305 303 259
HKGBC43	HKGBC43R	168	blastx.2	(AF106037) adipocyte-derived leucine aminopeptidase [Homo sapiens]	gb AAF07395.1 AF106037_1	92%	1	267
HLICD96	HLICD96R	169	blastx.2	leucine-rich alpha-2-glycoprotein - human	pir A03211 NBHU A2	88%	170	277
HLJBC28	HLJBC28RP00E	170	blastx.2	(AK000496) unnamed protein product [Homo sapiens]	dbj BAA91205.1	62%	552	262
HMKDT37	HMKDT37R	173	blastx.2	(AL110224) hypothetical protein [Homo sapiens]	emb CAB53682.1	93% 53%	13 99	108 143
HMPTG96	HMPTG96R P00E	174	blastx.2	(AF133299) lectin-like NK cell receptor LLT1 [Homo sapiens]	gb AAF22159.1 AF133299_1	97%	85	546
HMTBX02	HMTBX02R P00	175	blastx.2	(AF133299) lectin-like NK cell receptor LLT1 [Homo sapiens]	gb AAF22159.1 AF133299_1	100%	27	131
HMXAB55	HMXAB55R	176	blastx.2	interleukin-2 receptor	gb AAB59535.1	72%	10	117

HTGDN55	1106269	156	WUblastx. 64	ISOFORM BETA OF Q9UKF5	sp_vs Q9UKF5- 01 Q9UKF5	26%	1864	2232
H7TWD35	H7TWD35F	158	blastx.2	GPR10 [Homo sapiens]	gb AAC50504.1	26%	2491	2880
HBCMB46	HBCMB46R	159	blastx.2	platelet glycoprotein IIIa beta chain (version 2) - human 1	pir A60798 A60798	31%	1705	1896
HCUBO07	HCUBO07R	161	blastx.2	hypothetical proline-rich protein 1 - polychaete (Owenia fusiformis) (fragment)	pir A34043 A34043	35%	1387	1530
HDPWO71	HDPWO71R	162	blastx.2	(AF024636) STE20-like kinase 3 [Homo sapiens]	gb AAB82560.1	23%	1384	1851
						36%	2245	2352
						35%	337	429
						23%	220	348
						24%	697	951
						29%	2272	2436
						27%	2596	2802
						25%	682	915
						33%	415	513
						47%	53	361
						54%	469	501
						98%	49	249
						72%	290	376
						83%	3	203
						95%	1	141
						95%	1	141
						95%	1	141
						95%	1	141
						85%	2	142
						85%	2	142
						85%	2	142
						85%	2	142
						78%	2	241

HAFAQ92	1106214	154	WUblastx. 64	(Q9HBW9) ETL PROTEIN.	Q9HBW9	38%	2527	2604
HLHCR16	1106227	155	WUblastx. 64	(Q9ES77) POLYDOM PROTEIN PRECURSOR.	Q9ES77	75%	88	2898
						32%	133	2613
						30%	184	2457
						28%	187	2760
						31%	199	1920
						28%	748	1644
						30%	229	882
						35%	202	615
						27%	1882	2436
						27%	403	1029
						28%	1711	2262
						31%	2326	2859
						34%	2509	2901
						26%	241	720
						32%	322	690
						32%	2263	2877
						35%	1261	1539
						29%	1714	2211
						33%	1987	2316
						32%	589	864
						28%	2128	2421
						26%	412	1137
						24%	736	1596
						23%	1315	1929
						31%	217	516
						23%	1552	2013
						39%	1708	1941

HMWCU6 3	1105747	149	WUblastx. 64	(Q9HD37) C-TYPE LECTIN (FRAGMENT).	Q9HD37	100%	73	366
HWADF26	1105857	150	WUblastx. 64	(Q9NZS2) LECTIN-LIKE RECEPTOR F1 (ACTIVATING CORECEPTOR NKP80).	Q9NZS2	94%	442	612
HDABT82	1106030	151	WUblastx. 64	(Q9H2W1) CDA01 (MS4A6A-POLYMORPH) (MS4A6A PROTEIN).	Q9H2W1	92%	339	947
HDPMQ42	1106192	152	WUblastx. 64	type II lectin-like immunoreceptor - human	pir JC7608 JC7608	93%	545	1285
HHFHQ39	1106209	153	WUblastx. 64	(Q9UGM3) DMBT1 PROTOTYPE PRECURSOR.	Q9UGM3	89%	253	963
						88%	7	2067
						64%	4	2022
						63%	7	1974
						95%	2010	3200
						62%	7	1974
						50%	2010	2372
						62%	7	2022
						86%	4	1218
						51%	895	1653
						61%	1624	1974
						83%	7	1227
						84%	7	1218
						87%	517	1218
						67%	1624	1947
						79%	73	1212
						85%	886	1227
						44%	2527	2607
						38%	2527	2628
						38%	2527	2604

HETAQ37	1103950	142	WUblastx. 64	Wnt inhibitory factor-1 - human	pir A59180 A59180	90%	244	1380
HCENW09	1104186	143	WUblastx. 64	hypothetical protein DKFZp434C131.1 - human (fragment)	pir T17265 T17265	99%	59	673
HAMFW0 5	1104784	144	WUblastx. 64	(Q9D4G6) 4932434F09RIK PROTEIN.	Q9D4G6	71%	2	64
HKACI03	1104932	145	WUblastx. 64	(AAG28379) F11 receptor protein.	AAG28379	92%	217	978
HDTEJ03	1104936	146	WUblastx. 64	(Q9H7K2) FLJ00070 PROTEIN (FRAGMENT).	Q9H7K2	97%	704	2251
						31%	665	2203
						29%	686	2206
						28%	704	2245
						27%	530	1879
						32%	1328	2212
						29%	479	1282
						38%	1595	1921
						43%	2027	2242
						32%	1412	1678
						27%	1037	1489
						36%	2018	2215
						33%	2069	2245
						30%	1463	1621
H7TSA02	1105584	147	WUblastx. 64	pituitary adenylate cyclase- activating polypeptide receptor 1	pir S33449 S33449	87%	682	11
H7TSA19	1105585	148	WUblastx. 64	pituitary adenylate cyclase activating peptide receptor type I precursor - human	pir JN0902 JN0902	85%	506	6

HE2DL46	1102938	139	WUblastx. 64	HEMBB1000317, w	Q9BQ16	47%	2546	2274
				(Q9BQ16) SIMILAR TO TESTICAN 3.		48%	2783	2616
						23%	2576	2106
						36%	2465	2391
						24%	407	321
						95%	160	1347
HDPSJ02	1103186	140	WUblastx. 64	(Q9HIH6) DJ1009E24.1.1 (A NOVEL PROTEIN SIMILAR TO THE MOUSE SIALOADH	Q9HIH6	95%	730	924
						99%	2	730
						30%	2	730
						33%	733	909
						91%	915	1553
						35%	104	754
						37%	5	754
						34%	733	906
						38%	2	754
						39%	730	888
						34%	5	754
						32%	733	942
						44%	924	1343
						35%	912	1313
						39%	906	1334
						34%	924	1340
						39%	912	1337
						34%	5	754
						40%	939	1349
						29%	733	894
HE6FT04	1103911	141	WUblastx. 64	hypothetical protein DKFZp566D213.1 - human	pir T08724 T08724	93%	201	1451

							36%	1097	1195
HSIFQ16	1100483	135	WUblastx. 64	(O75427) LEUCIN RICH NEURONAL PROTEIN.	O75427		27%	79	1083
HWLDY64	1100528	136	WUblastx. 64	pregnancy-specific glycoprotein IIA form c precursor - human	pir H43354 H43354		89% 82% 100%	116 1468 84	1393 1941 125
HELBB20	1100919	137	WUblastx. 64	(Q9WUG5) SLIT2.	Q9WUG5		32% 25%	382 1073	161 477
							33% 33% 32% 26% 28% 30% 26% 30% 31% 34% 28% 27% 28% 28% 30% 32% 29% 26% 27% 28%	2069 1517 1550 1517 2012 2069 1694 1610 1610 1826 1898 2075 1418 1538 1820 1715 1910 1826 1595 1571	1515 987 984 618 1158 1593 1227 1152 1233 1413 1449 1593 975 1158 1446 1380 1521 1371 1233 1299
HETJP84	1102857	138	WUblastx. 64	(BAB55049) CDNA FLJ14438 fis, clone	BAB55049		98% 51%	2378 2714	165 2445

HWLED94	1099509	131	WUblastx. 64	(Q9HAU8) RNPEP-LIKE PROTEIN.	Q9HAU8	19% 18% 67% 95%	1722 1530 1291 143	1306 1297 551 1624
HKAJI55	1100188	132	WUblastx. 64	(Q9Y5P9) ENDOCYTTIC RECEPTOR ENDO180.	Q9Y5P9	100% 29% 29% 26% 23% 25%	9 9 48 75 36 36	728 725 734 734 734 626
HCEPF75.	1100345	133	WUblastx. 64	(Q9UGS3) DJ756G23.1 (NOVEL LEUCINE RICH PROTEIN) (FRAGMENT).	Q9UGS3	86% 29%	1470 1470	421 448
HRDFA94	1100389	134	WUblastx. 64	(Q9H1H6) DJ1009E24.1.1 (A NOVEL PROTEIN SIMILAR TO THE MOUSE SIALOADH	Q9H1H6	98% 43% 40% 30% 42% 36% 35% 32% 98% 34% 32% 31% 29% 23% 25%	1058 1079 1058 361 1073 1034 1025 1103 10 580 583 277 13 811 4	1243 1243 1240 1059 1243 1240 1231 1240 1077 1053 1065 1053 1122 1002 786

HDPUN70	1096317	124	WUblastx. 64	(Q9BXN2) DENDRITIC CELL-ASSOCIATED C- TYPE LECTIN-1.	Q9BXN2	91%	92	709
HHEPF57	1096422	125	WUblastx. 64	(Q9H3W5) HYPOTHETICAL 79.4 KDA PROTEIN.	Q9H3W5	93%	864	2987
HFKFH60	1096718	126	WUblastx. 64	(AAC51260) Matrilin-2 precursor.	AAC51260	95% 38% 41% 86%	495 495 229 79	2144 794 495 495
HHBFK81	1096722	127	WUblastx. 64	SHORT ISOFORM OF O00339	sp_vs O00339- 01 O00339	98% 58% 51% 50% 47% 47% 47% 40%	408 414 399 414 390 390 438 945	1781 944 944 944 944 944 944 1511
HARNY39	1097714	128	WUblastx. 64	(O75462) CYTOKINE- LIKE FACTOR-1 PRECURSOR.	O75462	85%	38	1192
HJPDH20	1099283	129	WUblastx. 64	(Q9H733) CDNA: FLJ21446 FTS, CLONE COL04458.	Q9H733	85% 84%	863 201	1090 845
HWLHB64	1099467	130	WUblastx. 64	(Q9PTD7) CINGULIN.	Q9PTD7	21% 34% 19% 22% 21%	1911 1899 1692 1713 1713	1414 1294 1294 1297 1210

HNECM88	1095640		64	(PROTEIN FOR MGC:4595).	O15389				
HCQBI77	1095827	114	WUblastx. 64	(O15389) OB BINDING PROTEIN-2 (SIGLEC5).	Q9CYA0	72%	4	573	
HBXFI23	1095877	115	WUblastx. 64	(Q9CYA0) 5730592L2IRIK PROTEIN.	sp_vs P06734-01 P06734	60% 70%	1141 965	938 267	
HEOAD12	1095888	116	WUblastx. 64	ISOFORM B OF P06734	Q9H1J4	39% 31%	798 220	866 639	
HEMFC27	1095935	117	WUblastx. 64	(Q9H1J4) HEDGEHOG-INTERACTING PROTEIN.	Q98SR5	75%	581	1075	
HFOYM21	1096023	118	WUblastx. 64	(Q98SR5) T CELL ANTIGEN CD44 ISOFORM B.	ITAH_HUMAN	37%	174	572	
HCE3H71	1096278	119	WUblastx. 64	(Q9UKX5) INTEGRIN ALPHA-11 PRECURSOR.	pir I52657 I52657	99% 90%	73 3173	3159 3637	
HOABR66	1096304	120	WUblastx. 64	seizure-related protein SEZ-6 precursor - mouse	Q9Y5P9	92% 37% 61% 46%	5 314 562 926	682 493 954 997	
HOHBI59	1096305	121	WUblastx. 64	(Q9Y5P9) ENDOCYTIC RECEPTOR ENDO180.	Q9Y5P9	87% 27% 28%	3 24 48	440 398 428	
HDPKI84	1096306	122	WUblastx. 64	(Q9Y5P9) ENDOCYTIC RECEPTOR ENDO180.	Q9BYU4	93% 24% 24% 22%	69 75 114 219	1523 1625 1547 1487	
		123	WUblastx. 64	(Q9BYU4) P2Y11 RECEPTOR.		42% 80%	712 881	885 943	

			64	ENDOTHELIAL MARKER 1 PRECURSOR (ENDOSIALIN PROTEIN).		33%	254 97	343 411
HISAP60	1094890	106	WUblastx. 64	(Q9BX67) JUNCTIONAL ADHESION MOLECULE 3 PRECURSOR.	Q9BX67	92%	540	1397
HWAGT47	1094891	107	WUblastx. 64	(Q9NZS2) LECTIN-LIKE RECEPTOR F1 (ACTIVATING CORECEPTOR NKP80).	Q9NZS2	78% 100%	4 174	174 608
HSXDF41	1094953	108	WUblastx. 64	(BAB55404) CDNA FLJ14935 fis, clone PLACE1009992, w	BAB55404	95% 99%	312 3	1655 323
HMTMD8 8	1095222	109	WUblastx. 64	(Q9NY25) MYELOID DAP12-ASSOCIATING LECTIN.	Q9NY25	92%	187	750
HEMEA55	1095289	110	WUblastx. 64	(Q75578) INTEGRIN ALPHA-10 PRECURSOR.	ITAG_HUMAN	92% 53%	125 736	802 828
HLJBH31	1095303	111	WUblastx. 64	(Q9Y286) QA79 MEMBRANE PROTEIN, ALLELIC VARIANT	Q9Y286	65% 65% 52%	575 173 1858	1876 643 1908
HFTCV04	1095439	112	WUblastx. 64	AIRM-1B PRECURSOR. (Q9BXN1) ASPARTATE- END LEUCINE-RICH REPEAT PROTEIN PRECURSOR.	Q9BXN1	70% 31% 29% 30% 92%	1049 736 535 529 283	1129 1044 939 999 1041
HMCHH31	1095550	113	WUblastx.	(Q9BR57) UNKNOWN	Q9BR57	87%	139	1194

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HMCHG54	1093752	94	WUblastx. 64	(BAB55362) CDNA FLJ14874 fis, clone PLACE1002722, w	BAB55362	33%	1183	1500
HELGM30	1093937	95	WUblastx. 64	(Q9PTE0) NEURONAL TETRASPANIN.	Q9PTE0	58%	640	840
HLKAB61	1094201	96	WUblastx. 64	(Q04721) NOTCH 2 PRECURSOR.	Q04721	80%	154	372
HFIZB56	1094512	97	WUblastx. 64	(BAB55420) CDNA FLJ14964 fis, clone PLACE4000581, m	BAB55420	100%	731	799
HAGFJ46	1094598	98	WUblastx. 64	(Q9HBW1) NAG14.	Q9HBW1	96%	2	1330
HYAAM4 3	1094663	99	WUblastx. 64	(Q9ULY5) MACROPHAGE C-TYPE LECTIN MINCLE.	Q9ULY5	100%	149	805
HDAAN31	1094668	100	WUblastx. 64	(Q9UKV4) COXSACKIE AND ADENOVIRUS RECEPTOR PROTEIN (FRAGMENT).	Q9UKV4	45%	708	806
HNGEO79	1094775	101	WUblastx. 64	Fc gamma (IgG) receptor high affinity - mouse	pir 48471 48471	46%	230	520
HWHQR2 5	1094777	102	WUblastx. 64	Delta-4 protein - human	pir JC7570 JC7570	80%	2	571
						40%	8	424

HEPNE19	1092413	88	WUblastx. 64	(Q9VZZ4) PXN PROTEIN.	Q9VZZ4	25% 39% 40% 37%	344 849 831 80	826 977 977 844
HLDAB75	1092493	89	WUblastx. 64	alpha-1-B-glycoprotein - human	pir A02113 OMHU 1B	46% 29% 37% 37% 30% 36% 29%	16 19 2845 232 232 765 1	3027 492 3069 495 507 854 282
HMSFP50	1092497	90	WUblastx. 64	Fc gamma (IgG) receptor high affinity - mouse	pir A46480 A46480	38%	78	914
HNTDM29	1092545	91	WUblastx. 64	CL1AA protein - rat	pir T17138 T17138	75% 74%	336 74	419 334
HATCZ07	1093241	92	WUblastx. 64	(Q9CRY3) 3100001108RIK PROTEIN (FRAGMENT).	Q9CRY3	94% 99%	13 140	63 1021
HDPXT07	1093488	93	WUblastx. 64	garp precursor - human	pir S42799 S42799	38% 42% 32% 29% 31% 26% 30% 33% 30% 31%	285 1153 1306 1147 1614 1159 1174 1306 1174 1162	965 1593 1641 1446 1898 1512 1536 1632 1494 1539

HTEEF19	1092204	84	WUblastx. 64	hypothetical protein DKFZp586B2420.1 - human (fragment)	pir T08774 T08774	30%	156	659
HEEAE73	1092290	85	WUblastx. 64	(O95137) CELL CYCLE RELATED KINASE (CYCLIN-DEPENDENT PROTEIN KINASE H	O95137	99%	251	745
H2CBE03	1092357	86	WUblastx. 64	(Q61846) MATERNAL EMBRYONIC LEUCINE ZIPPER KINASE (SERINE/THREONINE K	Q61846	92%	86	754
HMEBH77	1092363	87	WUblastx. 64	(BAB55399) CDNA FLJ14927 fis, clone PLACE1009094, w	BAB55399	92%	47	799
						35%	798	977
						31%	831	938
						38%	74	844
						31%	113	841
						43%	822	977

HAMFJ04	1091042	75	WUblastx. 64	(Q9D676) 4632413B12RIK PROTEIN.	Q9D676	59%	751	975
HAIDK30	1091076	76	WUblastx. 64	(Q9HBX2) AMINOPEPTIDASE.	Q9HBX2	97%	2	2155
HCEIR41	1091358	77	WUblastx. 64	(Q9H284) SEROLOGICALLY DEFINED BREAST CANCER ANTIGEN NY- BR-38.	Q9H284	37% 37% 34%	60 81 96	377 377 377
HCESF90	1091570	78	WUblastx. 64	(Q9NS59) ZINC FINGER PROTEIN (FRAGMENT).	Q9NS59	81% 99% 37% 33% 79%	3651 4561 4297 3661 4214	3142 4193 4211 3557 3684
HMSFT51	1091878	79	WUblastx. 64	(Q9UKQ0) TYPE II TRANSMEMBRANE PROTEIN MDL-1.	Q9UKQ0	100%	3	134
HAICQ70	1091887	80	WUblastx. 64	(Q9ET54) ACTIN- ASSOCIATED PROTEIN PALLADIN (FRAGMENT).	Q9ET54	89% 93% 36% 76%	267 1959 591 3231	1463 2105 869 3281
HDQGZ89	1092002	81	WUblastx. 64	(Q9H2N6) DC22.	Q9H2N6	69%	114	581
HWAFW3 9	1092095	82	WUblastx. 64	(Q9UKY2) ADIPOCYTE- DERIVED LEUCINE AMINOPEPTIDASE.	Q9UKY2	98%	200	367
HE8UL90	1092181	83	WUblastx. 64	(P82279) CRUMBS HOMOLOG 1.	P82279	80% 33% 29%	78 150 156	701 749 758

HBJHS02	1085743	65	64	WUblastx. 64	ALPHA-11 PRECURSOR. (Q9BR57) UNKNOWN (PROTEIN FOR MGC:4595).	Q9BR57	68% 45%	64 4	651 297
HE8PB92	1086828	66	64	WUblastx. 64	(Q9VYF1) CG15744 PROTEIN.	Q9VYF1	29% 28% 33%	65 532 467	355 669 538
HTEMU66	1086856	67	64	WUblastx. 64	(Q9H5T2) CDNA: FLJ23074 FIS, CLONE LNG05744.	Q9H5T2	98%	603	962
HE2PQ73	1089285	68	64	WUblastx. 64	(BAB55023) CDNA FLJ14391 fis, clone HEMBA1003077, w	BAB55023	53% 27% 56%	265 343 929	909 897 1174
HOHEC71	1090538	69	64	WUblastx. 64	(Q9Y5D7) PROTOCOLADHERIN GAMMA A12 SHORT FORM PROTEIN.	Q9Y5D7	86% 33%	66 90	617 506
HCE3J64	1090546	70	64	WUblastx. 64	(O60344) ENDOTHELIN- CONVERTING ENZYME 2 (EC 3.4.24.71) (ECE	ECE2_HUMAN	98%	3	647
HCE3L81	1090711	71	64	WUblastx. 64	(Q9BQH3) HYPOTHETICAL 55.7 KDA PROTEIN.	Q9BQH3	88% 97%	677 91	703 666
HUFDI62	1090753	72	64	WUblastx. 64	(O75296) R29124_1.	O75296	90% 99%	601 95	876 517
HE8FC96	1090865	73	64	WUblastx. 64	(Q9NYK3) PROTEIN KINASE MEKK2B.	Q9NYK3	80% 99%	108 947	965 1648
HTEPC92	1091037	74	64	WUblastx. 64	(Q9P126) C-TYPE LECTIN-LIKE RECEPTOR-2.	Q9P126	26% 78%	1101 386	1247 535

HPLBQ44	1081779	54	WUblastx. 64	(Q9H0U1) HYPOTHETICAL 30.7 KDA PROTEIN.	Q9H0U1	82%	177	986
HAHBE26	1082286	55	WUblastx. 64	(Q9EQJ5) STRIATED MUSCLE-SPECIFIC SERINE/THREONINE PROTEIN KINASE.	Q9EQJ5	35% 57% 42%	109 694 172	657 750 660
HMMCH0 4	1082413	56	WUblastx. 64	(Q9H387) PRO2550.	Q9H387	51% 61% 70%	683 724 591	585 686 499
HOFMN17	1083062	57	WUblastx. 64	CL1AA protein - rat	pir T17138 T17138	75%	15	791
HKABY22	1083735	58	WUblastx. 64	hypothetical protein DKFZp434K2323.1 - human (fragment)	pir T46418 T46418	74% 86%	702 271	1022 678
HLDQJ74	1083901	59	WUblastx. 64	(BAB55023) CDNA FLJ14391 fis, clone HEMBA1003077, w	BAB55023	92%	65	1918
HTLDS55	1083991	60	WUblastx. 64	(Q9D9U8) 1700028G04RIK PROTEIN.	Q9D9U8	82%	340	1089
HDQ GK06	1084465	61	WUblastx. 64	(Q9H013) MELTRIN- BETA/ADAM 19 HOMOLOGUE.	Q9H013	93%	366	1910
HNGME19	1084997	62	WUblastx. 64	(Q9NRT9) PROTOCOLADHERIN 10 (FRAGMENT).	Q9NRT9	96% 32% 27%	3 207 6	1187 857 881
HELGO48	1085552	63	WUblastx. 64	(O60687) SUSHI- REPEAT PROTEIN.	O60687	99%	28	1296
HRDAF83	1085723	64	WUblastx.	(Q9UKX5) INTEGRIN	TTAH_HUMAN	93%	992	696

			64	9030421L11RIK PROTEIN.					
HNFHW14	1057738	45	WUblastx. 64	(Q9QZT2) SERPENTINE RECEPTOR.	Q9QZT2		44%	38	256
HUKDS43	1061241	46	WUblastx. 64	(Q9ERK3) NEPRILYSIN- LIKE PEPTIDASE ALPHA.	Q9ERK3		88% 82%	8 420	313 755
HAOAD02	1063218	47	WUblastx. 64	(O75578) INTEGRIN ALPHA-10 PRECURSOR.	ITAG_HUMAN		86% 34% 97% 42%	444 4912 1 5013	3455 4995 540 5069
HOUCD83	1070724	48	WUblastx. 64	(Q9NQS3) NECTIN 3.	Q9NQS3		87%	3892	2978
HOFMS43	1071027	49	WUblastx. 64	(O02839) PORCINE MEMBRANE COFACTOR PROTEIN.	O02839		51% 29%	73 97	573 525
HMTAK58	1071812	50	WUblastx. 64	(Q9UHP7) LECTIN-LIKE NK CELL RECEPTOR LLT1 (C-TYPE LECTIN).	Q9UHP7		98%	2	553
H7TBB38	1073467	51	WUblastx. 64	(Q9NSD7) G-PROTEIN COUPLED RECEPTOR SALPR.	Q9NSD7		61% 52% 41%	78 543 330	233 692 467
HPWBS63	1077575	52	WUblastx. 64	(Q9EPL2) CALSYNTENIN-1 PROTEIN PRECURSOR.	Q9EPL2		91%	15	833
HSDJH63	1078285	53	WUblastx. 64	(Q9NYC3) TOLL-LIKE RECEPTOR 9 FORM A.	Q9NYC3		100% 52% 45% 58% 92%	107 128 122 131 192	190 184 181 181 1358

H7TSC95	997589	33	WUblastx. 64	pituitary adenylate cyclase activating peptide receptor type I precursor - human	pir JN0902 JN0902	86%	456	1
HTEIS02	999204	34	WUblastx. 64	(Q9NV82) CDNA FLJ10877 FIS, CLONE NT2RP4001861 (FRAGMENT).	Q9NV82	41% 100% 37% 100%	130 93 98 683	216 122 202 787
HTXQD18	1000599	35	WUblastx. 64	(Q9NW37) CDNA FLJ10337 FIS, CLONE NT2RM2000714, WEAKLY SIMILAR TO RAS	Q9NW37	96% 39%	2 116	553 535
HPMSA16	1011769	37	WUblastx. 64	(Q9H7M4) FLJ00046 PROTEIN (FRAGMENT).	Q9H7M4	92% 97%	166 294	47 163
HTTKE35	1027025	38	WUblastx. 64	(Q9DAW5) 1600015H20RIK PROTEIN.	Q9DAW5	90%	133	780
HCRPE25	1034245	39	WUblastx. 64	(Q9BY18) MS4A4A PROTEIN.	Q9BY18	93%	316	975
HWAGX4 9	1034246	40	WUblastx. 64	(Q9BY18) MS4A4A PROTEIN.	Q9BY18	94%	208	867
HTENW06	1040030	41	WUblastx. 64	(Q9VEK6) CG5407 PROTEIN.	Q9VEK6	34% 34%	300 312	785 677
HODFW19	1052332	42	WUblastx. 64	(Q9Y5D1) PROTOCADHERIN GAMMA A6 SHORT FORM PROTEIN.	Q9Y5D1	48% 32% 26% 90%	1239 5 5 2	1379 793 847 1249
HACAD35	1052372	43	WUblastx. 64	(Q9UDN0) SIMILAR TO COCH-5B2.	Q9UDN0	90%	4	597
HHPDX35	1056108	44	WUblastx.	(Q9D332)	Q9D332	72%	12	764

HHERQ79	985266	27	WUblastx. 64	(Q14164) INHIBITOR OF NUCLEAR FACTOR KAPPA-B KINASE EPSILON	IKKE_HUMAN	100%	361	711
HRABV49	988947	28	WUblastx. 64	(Q9H7K2) FLJ00070 PROTEIN (FRAGMENT).	Q9H7K2	94% 29% 29% 29% 39% 38% 43% 32% 37% 27% 30% 36% 33% 33% 30%	6 78 78 33 402 1092 261 909 138 534 78 1515 1515 1566 960	1748 1742 1703 1376 779 1418 479 1175 320 986 509 1712 1727 1742 1118
HNGGK47	993602	29	WUblastx. 64	(Q9XRX5) HHLA3 PROTEIN (FRAGMENT).	Q9XRX5	76%	277	366
HAJRC82	993759	30	WUblastx. 64	(Q9H075) HYPOTHETICAL 81.8 KDA PROTEIN.	Q9H075	55% 25%	44 1139	2005 1960
HJABW64	994001	31	WUblastx. 64	(Q9HCZ4) DJ677HI5.1 (A NOVEL PROTEIN SIMILAR TO LEUCINE- RICH REPEAT P	Q9HCZ4	99%	25	456
HAJBT01	997082	32	WUblastx. 64	nucleophosmin - human	pir A33423 A32915	74% 74%	1174 1293	1290 2012

HBKDR69	956894	23	WUblastx. 64	(Q9UBF9) MYOTILIN.	Q9UBF9		32%	5	865
HSIGI60	956986	24	WUblastx. 64	dipeptidyl aminopeptidase-like protein (EC 3.4.14.-) short form, DPPX-S - bovine	pir A41793 A41793		48%	1218	865 835 850 856 706 865 903 835 868 897 868 439 898 169 868 874 865
HOFNI12	982193	25	WUblastx. 64	(Q9Y3E9) CGI-151 PROTEIN.	Q9Y3E9		84%	18	485
HEGAZ61	983192	26	WUblastx. 64	(Q9H2U9) A DISINTEGRIN AND METALLOPROTEINASE 7.	Q9H2U9		98%	60	560

				64	APYRASE-LIKE PROTEIN I. (Q9Y3Q7) TMDC III PROTEIN.	Q9Y3Q7							
HTEBO72	947922	19		WUblastx. 64					92%	30			554
HWLHQ59	949724	20		WUblastx. 64	(Q9NXJ0) CDNA FLJ20217 FIS, CLONE COLF3334.	Q9NXJ0			93%	1181			381
HDQES11	950838	21		WUblastx. 64	(Q9HBM5) DIPEPTIDYL PEPTIDASE 8.	Q9HBM5			97%	2030			282
HOHCCE96	955622	22		WUblastx. 64	(Q9ES77) POLYDOM PROTEIN PRECURSOR.	Q9ES77			63%	838			903
									66%	868			903
									45%	832			900
									31%	844			900
									35%	853			903
									46%	856			900
									40%	847			903
									63%	868			900
									30%	2			568
									23%	2			496
									33%	539			868
									26%	56			373
									26%	125			751
									32%	2			835
									26%	14			559
									21%	146			529
									31%	8			160
									30%	2			751
									26%	5			868
									32%	2			844
									25%	5			817

H7TBE17	858079	14	WUblastx. 64	(Q9NSD7) G-PROTEIN COUPLED RECEPTOR SALPR.	Q9NSD7	26%	1094	1333 1112 1061 1028 604 1411 604 1133 1046 1070 1118 1330 619 2112 604 1974 2316 1755
H7TRA52	858086	15	WUblastx. 64	(Q9NSD7) G-PROTEIN COUPLED RECEPTOR SALPR.	Q9NSD7	56%	190	5 500 245
HSSJC65	872582	16	WUblastx. 64	(Q9BZR6) NOGO RECEPTOR.	Q9BZR6	90%	39	1361
HKLAA30	875001	17	WUblastx. 64	(Q9BWP8) UNKNOWN (PROTEIN FOR MGC:3279).	Q9BWP8	100%	2 103	94 222
HPJCP59	943763	18	WUblastx.	(Q9NOZ7) LYSOSOMAL	Q9NOZ7	91%	220	1026

TABLE 2

Clone ID NO:Z	Contig ID:	SEQ ID NO:X	Analysis Method	PFam/NR Description	PFam/NR Accession Number	Score/ Percent Identity	NT From	NT To
HEMEC34	684296	11	WUblastx. 64	(O95150) VASCULAR ENDOTHELIAL CELL GROWTH INHIBITOR.	O95150	100%	427	879
HDPMQ69	812744	12	WUblastx. 64	(Q9NYZ4) SIGLEC SAF2.	Q9NYZ4	64%	1188	1451
						57%	166	486
						34%	1736	1906
						50%	3168	3209
						33%	714	803
						41%	576	980
HTPAU91	846654	13	WUblastx. 64	(Q9H4W4) DJ947L8.1.8 (NOVEL SUSHI (SCR REPEAT) DOMAIN PROTEIN) (FRAGM	Q9H4W4	81%	657	905
						100%	604	693
						59%	735	1052
						97%	311	607
						31%	750	1124
						35%	1091	1222
						31%	1103	1219
						40%	628	705
						31%	1444	2112
						37%	323	499
						31%	1127	1255
						42%	631	693
						34%	1582	2154
						36%	1124	1342
						41%	1100	1219

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matches to EST and cDNA sequences contained in the NCBI (National Center for Biotechnology Information) UniGene database. Each sequence in the UniGene database is assigned to a "cluster"; all of the ESTs, cDNAs, and STSs in a cluster are believed to be derived from a single gene. Chromosomal mapping data is often available for one or more sequence(s) in a UniGene cluster; this data (if consistent) is then applied to the cluster as a whole. Thus, it is possible to infer the chromosomal location of a new polynucleotide sequence by determining its identity with a mapped UniGene cluster.

[0068] A modified version of the computer program BLASTN (Altschul et al., J. Mol. Biol. 215:403-410 (1990); and Gish and States, Nat. Genet. 3:266-272 (1993)) was used to search the UniGene database for EST or cDNA sequences that contain exact or near-exact matches to a polynucleotide sequence of the invention (the 'Query'). A sequence from the UniGene database (the 'Subject') was said to be an exact match if it contained a segment of 50 nucleotides in length such that 48 of those nucleotides were in the same order as found in the Query sequence. If all of the matches that met this criteria were in the same UniGene cluster, and mapping data was available for this cluster, it is indicated in Table 1 under the heading "Cytologic Band". Where a cluster had been further localized to a distinct cytologic band, that band is disclosed; where no banding information was available, but the gene had been localized to a single chromosome, the chromosome is disclosed.

[0069] Once a presumptive chromosomal location was determined for a polynucleotide of the invention, an associated disease locus was identified by comparison with a database of diseases which have been experimentally associated with genetic loci. The database used was the Morbid Map, derived from OMIM™ (*supra*). If the putative chromosomal location of a polynucleotide of the invention (Query sequence) was associated with a disease in the Morbid Map database, an OMIM reference identification number was noted in column 10, Table 1, labelled "OMIM Disease Reference(s)". Table 5 is a key to the OMIM reference identification numbers (column 1), and provides a description of the associated disease in Column 2.

1. It will be appreciated that depending on the analytical criteria used to predict antigenic determinants, the exact address of the determinant may vary slightly.

[0066] Column 8 in Table 1 provides an expression profile and library code: count for each of the contig sequences (SEQ ID NO:X) disclosed in Table 1, which can routinely be combined with the information provided in Table 4 and used to determine the tissues, cells, and/or cell line libraries which predominantly express the polynucleotides of the invention. The first number in column 8 (preceding the colon), represents the tissue/cell source identifier code corresponding to the code and description provided in Table 4. For those identifier codes in which the first two letters are not "AR", the second number in column 8 (following the colon) represents the number of times a sequence corresponding to the reference polynucleotide sequence was identified in the tissue/cell source. Those tissue/cell source identifier codes in which the first two letters are "AR" designate information generated using DNA array technology. Utilizing this technology, cDNAs were amplified by PCR and then transferred, in duplicate, onto the array. Gene expression was assayed through hybridization of first strand cDNA probes to the DNA array. cDNA probes were generated from total RNA extracted from a variety of different tissues and cell lines. Probe synthesis was performed in the presence of ^{33}P dCTP, using oligo(dT) to prime reverse transcription. After hybridization, high stringency washing conditions were employed to remove non-specific hybrids from the array. The remaining signal, emanating from each gene target, was measured using a Phosphorimager. Gene expression was reported as Phosphor Stimulating Luminescence (PSL) which reflects the level of phosphor signal generated from the probe hybridized to each of the gene targets represented on the array. A local background signal subtraction was performed before the total signal generated from each array was used to normalize gene expression between the different hybridizations. The value presented after "[array code]:" represents the mean of the duplicate values, following background subtraction and probe normalization. One of skill in the art could routinely use this information to identify normal and/or diseased tissue(s) which show a predominant expression pattern of the corresponding polynucleotide of the invention or to identify polynucleotides which show predominant and/or specific tissue and/or cell expression.

[0067] Column 9 in Table 1 provides a chromosomal map location for certain polynucleotides of the invention. Chromosomal location was determined by finding exact

[0062] The first column in Table 1 provides the gene number in the application corresponding to the clone identifier. The second column in Table 1 provides a unique "Clone ID NO:Z" for a cDNA clone related to each contig sequence disclosed in Table 1. This clone ID references the cDNA clone which contains at least the 5' most sequence of the assembled contig and at least a portion of SEQ ID NO:X was determined by directly sequencing the referenced clone. The reference clone may have more sequence than described in the sequence listing or the clone may have less. In the vast majority of cases, however, the clone is believed to encode a full-length polypeptide. In the case where a clone is not full-length, a full-length cDNA can be obtained by methods described elsewhere herein.

[0063] The third column in Table 1 provides a unique "Contig ID" identification for each contig sequence. The fourth column provides the "SEQ ID NO:" identifier for each of the contig polynucleotide sequences disclosed in Table 1. The fifth column, "ORF (From-To)", provides the location (i.e., nucleotide position numbers) within the polynucleotide sequence "SEQ ID NO:X" that delineate the preferred open reading frame (ORF) shown in the sequence listing and referenced in Table 1, column 6, as SEQ ID NO:Y. Where the nucleotide position number "To" is lower than the nucleotide position number "From", the preferred ORF is the reverse complement of the referenced polynucleotide sequence.

[0064] The sixth column in Table 1 provides the corresponding SEQ ID NO:Y for the polypeptide sequence encoded by the preferred ORF delineated in column 5. In one embodiment, the invention provides an amino acid sequence comprising, or alternatively consisting of, a polypeptide encoded by the portion of SEQ ID NO:X delineated by "ORF (From-To)". Also provided are polynucleotides encoding such amino acid sequences and the complementary strand thereto.

[0065] Column 7 in Table 1 lists residues comprising epitopes contained in the polypeptides encoded by the preferred ORF (SEQ ID NO:Y), as predicted using the algorithm of Jameson and Wolf, (1988) Comp. Appl. Biosci. 4:181-186. The Jameson-Wolf antigenic analysis was performed using the computer program PROTEAN (Version 3.11 for the Power MacIntosh, DNASTAR, Inc., 1228 South Park Street Madison, WI). In specific embodiments, polypeptides of the invention comprise, or alternatively consist of, at least one, two, three, four, five or more of the predicted epitopes as described in Table

						41, AR104: 41, AR182: 41, AR060: 39, AR286: 37, AR033: 36, AR277: 36, AR237: 35, AR061: 34, AR229: 33, AR226: 30, AR052: 29, AR313: 28, AR238: 26, AR232: 25, AR244: 23, AR039: 23, AR312: 19, AR265: 19, AR053: 17, AR310: 16, AR205: 14, AR263: 13, AR194: 13, AR213: 13, AR233: 12, AR246: 11, AR206: 11, AR204: 10, AR227: 7, AR202: 7
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176	HYBAU21	HYBAU21 RA	186	3 - 203	362	AR218: 222, AR219: 181, AR175: 136, AR256: 132, AR240: 127, AR253: 125, AR291: 117, AR273: 112, AR275: 108, AR179: 106, AR268: 102, AR290: 93, AR295: 89, AR248: 88, AR177: 83, AR314: 83, AR315: 79, AR292: 79, AR251: 78, AR280: 76, AR247: 75, AR269: 74, AR267: 73, AR185: 72, AR183: 71, AR296: 70, AR271: 70, AR055: 68, AR096: 65, AR089: 64, AR298: 64, AR259: 62, AR266: 61, AR285: 60, AR270: 59, AR192: 57, AR231: 57, AR274: 57, AR284: 56, AR234: 56, AR184: 55, AR316: 54, AR198: 54, AR186: 53, AR300: 51, AR282: 51, AR241: 50, AR258: 49, AR281: 49, AR243: 49, AR293: 48, AR294: 48, AR249: 48, AR289: 47, AR299: 42, AR283: 42, AR309:		
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175	HWCAA29	HWCAA29 R	185	1 - 417	361	AR259: 5, AR229: 5, AR300: 5, AR234: 4, AR263: 4, AR246: 4, AR060: 4, AR258: 4, AR039: 4, AR053: 4, AR226: 4, AR244: 4, AR205: 3, AR232: 3, AR204: 3, AR277: 3, AR206: 2, AR194: 2, AR233: 2, AR055: 1, AR283: 1, AR061: 1, AR281: 1, AR227: 1, AR202: 0			
						AR299: 372, AR290: 271, AR300: 239, AR268: 237, AR267: 214, AR270: 147, AR183: 135, AR234: 130, AR096: 114, AR213: 81, AR313: 78, AR231: 62, AR247: 60, AR240: 59, AR053: 48, AR282: 45, AR316: 37, AR238: 35, AR246: 35, AR249: 28, AR269: 26, AR055: 24, AR052: 24, AR289: 22, AR202: 20, AR256: 19, AR198: 18, AR241: 17, AR237: 14,			

[0908] Therapeutics of the invention are also suitably administered by sustained-release systems. Suitable examples of sustained-release Therapeutics are administered orally, rectally, parenterally, intracisternally, intravaginally, intraperitoneally, topically (as by powders, ointments, gels, drops or transdermal patch), buccally, or as an oral or nasal spray. "Pharmaceutically acceptable carrier" refers to a non-toxic solid, semisolid or liquid filler, diluent, encapsulating material or formulation auxiliary of any type. The term "parenteral" as used herein refers to modes of administration which include intravenous, intramuscular, intraperitoneal, intrasternal, subcutaneous and intraarticular injection and infusion.

[0909] Therapeutics of the invention are also suitably administered by sustained-release systems. Suitable examples of sustained-release Therapeutics include suitable polymeric materials (such as, for example, semi-permeable polymer matrices in the form of shaped articles, e.g., films, or microcapsules), suitable hydrophobic materials (for example as an emulsion in an acceptable oil) or ion exchange resins, and sparingly soluble derivatives (such as, for example, a sparingly soluble salt).

[0910] Sustained-release matrices include polylactides (U.S. Pat. No. 3,773,919, EP 58,481), copolymers of L-glutamic acid and gamma-ethyl-L-glutamate (Sidman et al., *Biopolymers* 22:547-556 (1983)), poly (2- hydroxyethyl methacrylate) (Langer et al., *J. Biomed. Mater. Res.* 15:167-277 (1981), and Langer, *Chem. Tech.* 12:98-105 (1982)), ethylene vinyl acetate (Langer et al., *Id.*) or poly-D- (-)-3-hydroxybutyric acid (EP 133,988).

[0911] Sustained-release Therapeutics also include liposomally entrapped Therapeutics of the invention (*see* generally, Langer, *Science* 249:1527-1533 (1990); Treat et al., in *Liposomes in the Therapy of Infectious Disease and Cancer*, Lopez-Berestein and Fidler (eds.), Liss, New York, pp. 317 -327 and 353-365 (1989)). Liposomes containing the Therapeutic are prepared by methods known per se: DE 3,218,121; Epstein et al., *Proc. Natl. Acad. Sci. (USA)* 82:3688-3692 (1985); Hwang et al., *Proc. Natl. Acad. Sci.(USA)* 77:4030-4034 (1980); EP 52,322; EP 36,676; EP 88,046; EP 143,949; EP 142,641; Japanese Pat. Appl. 83-118008; U.S. Pat. Nos. 4,485,045 and 4,544,545; and EP 102,324. Ordinarily, the liposomes are of the small (about 200-800 Angstroms) unilamellar type in which the lipid content is greater than about 30 mol. percent cholesterol, the selected proportion being adjusted for the optimal Therapeutic.

Example 13: Formulation

[0904] The invention also provides methods of treatment and/or prevention of diseases or disorders (such as, for example, any one or more of the diseases or disorders disclosed herein) by administration to a subject of an effective amount of a Therapeutic. By therapeutic is meant polynucleotides or polypeptides of the invention (including fragments and variants), agonists or antagonists thereof, and/or antibodies thereto, in combination with a pharmaceutically acceptable carrier type (e.g., a sterile carrier).

[0905] The Therapeutic will be formulated and dosed in a fashion consistent with good medical practice, taking into account the clinical condition of the individual patient (especially the side effects of treatment with the Therapeutic alone), the site of delivery, the method of administration, the scheduling of administration, and other factors known to practitioners. The "effective amount" for purposes herein is thus determined by such considerations.

[0906] As a general proposition, the total pharmaceutically effective amount of the Therapeutic administered parenterally per dose will be in the range of about 1ug/kg/day to 10 mg/kg/day of patient body weight, although, as noted above, this will be subject to therapeutic discretion. More preferably, this dose is at least 0.01 mg/kg/day, and most preferably for humans between about 0.01 and 1 mg/kg/day for the hormone. If given continuously, the Therapeutic is typically administered at a dose rate of about 1 ug/kg/hour to about 50 ug/kg/hour, either by 1-4 injections per day or by continuous subcutaneous infusions, for example, using a mini-pump. An intravenous bag solution may also be employed. The length of treatment needed to observe changes and the interval following treatment for responses to occur appears to vary depending on the desired effect.

[0907] Therapeutics can be are administered orally, rectally, parenterally, intracisternally, intravaginally, intraperitoneally, topically (as by powders, ointments, gels, drops or transdermal patch), buccally, or as an oral or nasal spray. "Pharmaceutically acceptable carrier" refers to a non-toxic solid, semisolid or liquid filler, diluent, encapsulating material or formulation auxiliary of any. The term "parenteral" as used herein refers to modes of administration which include intravenous, intramuscular, intraperitoneal, intrasternal, subcutaneous and intraarticular injection and infusion.

Example 12: Method of Detecting Abnormal Levels of a Polypeptide in a Biological Sample

[0899] A polypeptide of the present invention can be detected in a biological sample, and if an increased or decreased level of the polypeptide is detected, this polypeptide is a marker for a particular phenotype. Methods of detection are numerous, and thus, it is understood that one skilled in the art can modify the following assay to fit their particular needs.

[0900] For example, antibody-sandwich ELISAs are used to detect polypeptides in a sample, preferably a biological sample. Wells of a microtiter plate are coated with specific antibodies, at a final concentration of 0.2 to 10 ug/ml. The antibodies are either monoclonal or polyclonal and are produced by the method described in Example 10. The wells are blocked so that non-specific binding of the polypeptide to the well is reduced.

[0901] The coated wells are then incubated for > 2 hours at RT with a sample containing the polypeptide. Preferably, serial dilutions of the sample should be used to validate results. The plates are then washed three times with deionized or distilled water to remove unbound polypeptide.

[0902] Next, 50 ul of specific antibody-alkaline phosphatase conjugate, at a concentration of 25-400 ng, is added and incubated for 2 hours at room temperature. The plates are again washed three times with deionized or distilled water to remove unbound conjugate.

[0903] Add 75 ul of 4-methylumbelliferyl phosphate (MUP) or p-nitrophenyl phosphate (NPP) substrate solution to each well and incubate 1 hour at room temperature. Measure the reaction by a microtiter plate reader. Prepare a standard curve, using serial dilutions of a control sample, and plot polypeptide concentration on the X-axis (log scale) and fluorescence or absorbance of the Y-axis (linear scale). Interpolate the concentration of the polypeptide in the sample using the standard curve.

Suggested PCR conditions consist of 35 cycles at 95 degrees C for 30 seconds; 60-120 seconds at 52-58 degrees C; and 60-120 seconds at 70 degrees C, using buffer solutions described in Sidransky et al., Science 252:706 (1991).

[0895] PCR products are then sequenced using primers labeled at their 5' end with T4 polynucleotide kinase, employing SequiTherm Polymerase (Epicentre Technologies). The intron-exon boundaries of selected exons is also determined and genomic PCR products analyzed to confirm the results. PCR products harboring suspected mutations are then cloned and sequenced to validate the results of the direct sequencing.

[0896] PCR products are cloned into T-tailed vectors as described in Holton et al., Nucleic Acids Research, 19:1156 (1991) and sequenced with T7 polymerase (United States Biochemical). Affected individuals are identified by mutations not present in unaffected individuals.

[0897] Genomic rearrangements are also observed as a method of determining alterations in a gene corresponding to a polynucleotide. Genomic clones isolated according to Example 2 are nick-translated with digoxigenin deoxy-uridine 5'-triphosphate (Boehringer Mannheim), and FISH performed as described in Johnson et al., Methods Cell Biol. 35:73-99 (1991). Hybridization with the labeled probe is carried out using a vast excess of human cot-1 DNA for specific hybridization to the corresponding genomic locus.

[0898] Chromosomes are counterstained with 4,6-diamino-2-phenylidole and propidium iodide, producing a combination of C- and R-bands. Aligned images for precise mapping are obtained using a triple-band filter set (Chroma Technology, Brattleboro, VT) in combination with a cooled charge-coupled device camera (Photometrics, Tucson, AZ) and variable excitation wavelength filters. (Johnson et al., Genet. Anal. Tech. Appl., 8:75 (1991)). Image collection, analysis and chromosomal fractional length measurements are performed using the ISee Graphical Program System. (Inovision Corporation, Durham, NC.) Chromosome alterations of the genomic region hybridized by the probe are identified as insertions, deletions, and translocations. These alterations are used as a diagnostic marker for an associated disease.

blocked with 2% Marvel-PBS for 2 hours at 37°C and then washed 3 times in PBS. Approximately 10^{13} TU of phage is applied to the tube and incubated for 30 minutes at room temperature tumbling on an over and under turntable and then left to stand for another 1.5 hours. Tubes are washed 10 times with PBS 0.1% Tween-20 and 10 times with PBS. Phage are eluted by adding 1 ml of 100 mM triethylamine and rotating 15 minutes on an under and over turntable after which the solution is immediately neutralized with 0.5 ml of 1.0M Tris-HCl, pH 7.4. Phage are then used to infect 10 ml of mid-log E. coli TG1 by incubating eluted phage with bacteria for 30 minutes at 37°C. The E. coli are then plated on TYE plates containing 1% glucose and 100 µg/ml ampicillin. The resulting bacterial library is then rescued with delta gene 3 helper phage as described above to prepare phage for a subsequent round of selection. This process is then repeated for a total of 4 rounds of affinity purification with tube-washing increased to 20 times with PBS, 0.1% Tween-20 and 20 times with PBS for rounds 3 and 4.

[0893] *Characterization of Binders.* Eluted phage from the 3rd and 4th rounds of selection are used to infect E. coli HB 2151 and soluble scFv is produced (Marks, et al., 1991) from single colonies for assay. ELISAs are performed with microtitre plates coated with either 10 pg/ml of the polypeptide of the present invention in 50 mM bicarbonate pH 9.6. Clones positive in ELISA are further characterized by PCR fingerprinting (see, e.g., International Publication No. WO 92/01047) and then by sequencing. These ELISA positive clones may also be further characterized by techniques known in the art, such as, for example, epitope mapping, binding affinity, receptor signal transduction, ability to block or competitively inhibit antibody/antigen binding, and competitive agonistic or antagonistic activity.

Example 11: Method of Determining Alterations in a Gene Corresponding to a Polynucleotide

[0894] RNA isolated from entire families or individual patients presenting with a phenotype of interest (such as a disease) is isolated. cDNA is then generated from these RNA samples using protocols known in the art. (See, Sambrook.) The cDNA is then used as a template for PCR, employing primers surrounding regions of interest in SEQ ID NO:X; and/or the nucleotide sequence of the cDNA contained in Clone ID NO:Z.

Isolation Of Antibody Fragments Directed Against A Polypeptide of the Invention From A Library Of scFvs

[0889] Naturally occurring V-genes isolated from human PBLs are constructed into a library of antibody fragments which contain reactivities against a polypeptide of the invention to which the donor may or may not have been exposed (see e.g., U.S. Patent 5,885,793 incorporated herein by reference in its entirety).

[0890] *Rescue of the Library.* A library of scFvs is constructed from the RNA of human PBLs as described in International Publication No. WO 92/01047. To rescue phage displaying antibody fragments, approximately 10^9 *E. coli* harboring the phagemid are used to inoculate 50 ml of 2xTY containing 1% glucose and 100 μ g/ml of ampicillin (2xTY-AMP-GLU) and grown to an O.D. of 0.8 with shaking. Five ml of this culture is used to inoculate 50 ml of 2xTY-AMP-GLU, 2×10^8 TU of delta gene 3 helper (M13 delta gene III, see International Publication No. WO 92/01047) are added and the culture incubated at 37°C for 45 minutes without shaking and then at 37°C for 45 minutes with shaking. The culture is centrifuged at 4000 r.p.m. for 10 min. and the pellet resuspended in 2 liters of 2xTY containing 100 μ g/ml ampicillin and 50 μ g/ml kanamycin and grown overnight. Phage are prepared as described in International Publication No. WO 92/01047.

[0891] M13 delta gene III is prepared as follows: M13 delta gene III helper phage does not encode gene III protein, hence the phage(mid) displaying antibody fragments have a greater avidity of binding to antigen. Infectious M13 delta gene III particles are made by growing the helper phage in cells harboring a pUC19 derivative supplying the wild type gene III protein during phage morphogenesis. The culture is incubated for 1 hour at 37°C without shaking and then for a further hour at 37°C with shaking. Cells are spun down (IEC-Centra 8,400 r.p.m. for 10 min), resuspended in 300 ml 2xTY broth containing 100 μ g ampicillin/ml and 25 μ g kanamycin/ml (2xTY-AMP-KAN) and grown overnight, shaking at 37°C. Phage particles are purified and concentrated from the culture medium by two PEG-precipitations (Sambrook et al., 1990), resuspended in 2 ml PBS and passed through a 0.45 μ m filter (Minisart NML; Sartorius) to give a final concentration of approximately 10^{13} transducing units/ml (ampicillin-resistant clones).

[0892] *Panning of the Library.* Immunotubes (Nunc) are coated overnight in PBS with 4 ml of either 100 μ g/ml or 10 μ g/ml of a polypeptide of the present invention. Tubes are

serum (inactivated at about 56°C), and supplemented with about 10 g/l of nonessential amino acids, about 1,000 U/ml of penicillin, and about 100 µg/ml of streptomycin.

[0886] The splenocytes of such mice are extracted and fused with a suitable myeloma cell line. Any suitable myeloma cell line may be employed in accordance with the present invention; however, it is preferable to employ the parent myeloma cell line (SP2O), available from the ATCC. After fusion, the resulting hybridoma cells are selectively maintained in HAT medium, and then cloned by limiting dilution as described by Wands et al. (Gastroenterology 80:225-232 (1981)). The hybridoma cells obtained through such a selection are then assayed to identify clones which secrete antibodies capable of binding the polypeptide of the invention.

[0887] Alternatively, additional antibodies capable of binding to a polypeptide of the invention can be produced in a two-step procedure using anti-idiotypic antibodies. Such a method makes use of the fact that antibodies are themselves antigens, and therefore, it is possible to obtain an antibody which binds to a second antibody. In accordance with this method, protein specific antibodies are used to immunize an animal, preferably a mouse. The splenocytes of such an animal are then used to produce hybridoma cells, and the hybridoma cells are screened to identify clones which produce an antibody whose ability to bind to the polypeptide of the invention-specific antibody can be blocked by the polypeptide of the invention. Such antibodies comprise anti-idiotypic antibodies to the polypeptide of the invention-specific antibody and are used to immunize an animal to induce formation of further polypeptide of the invention-specific antibodies.

[0888] For *in vivo* use of antibodies in humans, an antibody is "humanized". Such antibodies can be produced using genetic constructs derived from hybridoma cells producing the monoclonal antibodies described above. Methods for producing chimeric and humanized antibodies are known in the art and are discussed herein. (See, for review, Morrison, Science 229:1202 (1985); Oi et al., BioTechniques 4:214 (1986); Cabilly et al., U.S. Patent No. 4,816,567; Taniguchi et al., EP 171496; Morrison et al., EP 173494; Neuberger et al., WO 8601533; Robinson et al., International Publication No. WO 8702671; Boulianne et al., Nature 312:643 (1984); Neuberger et al., Nature 314:268 (1985)).

GCCGCGGGAGGAGCAGTACAACAGCACGTACCGTGTGGTCAGCGTC
 CTCACCGTCCTGCACCAGGACTGGCTGAATGGCAAGGAGTACAAGT
 GCAAGGTCTCCAACAAAGCCCTCCCAACCCCCATCGAGAAAACCAT
 CTCCAAAGCCAAAGGGCAGCCCCGAGAACCACAGGTGTACACCCTG
 CCCCCATCCCGGGATGAGCTGACCAAGAACCAGGTCAGCCTGACCT
 GCCTGGTCAAAGGCTTCTATCCAAGCGACATCGCCGTGGAGTGGGA
 GAGCAATGGGCAGCCGGAGAACAACACTACAAGACCACGCCTCCCGTG
 CTGGACTCCGACGGCTCCTTCTTCCTCTACAGCAAGCTCACCGTGGA
 CAAGAGCAGGTGGCAGCAGGGGAACGTCTTCTCATGCTCCGTGATG
 CATGAGGCTCTGCACAACCACTACACGCAGAAGAGCCTCTCCCTGTC
 TCCGGGTAAATGAGTGCGACGGCCGCGACTCTAGAGGAT (SEQ ID
 NO: 1)

Example 10: Production of an Antibody from a Polypeptide

Hybridoma Technology

[0884] The antibodies of the present invention can be prepared by a variety of methods. (See, Current Protocols, Chapter 2.) As one example of such methods, cells expressing a polypeptide of the invention are administered to an animal to induce the production of sera containing polyclonal antibodies. In a preferred method, a preparation of a polypeptide of the invention is prepared and purified to render it substantially free of natural contaminants. Such a preparation is then introduced into an animal in order to produce polyclonal antisera of greater specific activity.

[0885] Monoclonal antibodies specific for a polypeptide of the invention are prepared using hybridoma technology (Kohler et al., Nature 256:495 (1975); Kohler et al., Eur. J. Immunol. 6:511 (1976); Kohler et al., Eur. J. Immunol. 6:292 (1976); Hammerling et al., in: Monoclonal Antibodies and T-Cell Hybridomas, Elsevier, N.Y., pp. 563-681 (1981)). In general, an animal (preferably a mouse) is immunized with a polypeptide of the invention or, more preferably, with a secreted polypeptide of the invention-expressing cell. Such polypeptide-expressing cells are cultured in any suitable tissue culture medium, preferably in Earle's modified Eagle's medium supplemented with 10% fetal bovine

protein facilitates purification. (See Example 5; see also EP A 394,827; Traunecker, et al., Nature 331:84-86 (1988)). Similarly, fusion to IgG-1, IgG-3, and albumin increases the half-life time *in vivo*. Nuclear localization signals fused to the polypeptides of the present invention can target the protein to a specific subcellular localization, while covalent heterodimer or homodimers can increase or decrease the activity of a fusion protein. Fusion proteins can also create chimeric molecules having more than one function. Finally, fusion proteins can increase solubility and/or stability of the fused protein compared to the non-fused protein. All of the types of fusion proteins described above can be made by modifying the following protocol, which outlines the fusion of a polypeptide to an IgG molecule, or the protocol described in Example 5.

[0881] Briefly, the human Fc portion of the IgG molecule can be PCR amplified, using primers that span the 5' and 3' ends of the sequence described below. These primers also should have convenient restriction enzyme sites that will facilitate cloning into an expression vector, preferably a mammalian expression vector.

[0882] For example, if pC4 (ATCC Accession No. 209646) is used, the human Fc portion can be ligated into the BamHI cloning site. Note that the 3' BamHI site should be destroyed. Next, the vector containing the human Fc portion is re-restricted with BamHI, linearizing the vector, and a polynucleotide of the present invention, isolated by the PCR protocol described in Example 1, is ligated into this BamHI site. Note that the polynucleotide is cloned without a stop codon, otherwise a fusion protein will not be produced.

[0883] If the naturally occurring signal sequence is used to produce the polypeptide of the present invention, pC4 does not need a second signal peptide. Alternatively, if the naturally occurring signal sequence is not used, the vector can be modified to include a heterologous signal sequence. (See, e.g., International Publication No. WO 96/34891.)

Human IgG Fc region:

```
GGGATCCGGAGCCCAAATCTTCTGACAAACTCACACATGCCACG
GTGCCCAGCACCTGAATTCGAGGGTGCACCGTCAGTCTTCCTCTTCC
CCCCAAAACCCAAGGACACCCTCATGATCTCCCGGACTCCTGAGGTC
ACATGCGTGGTGGTGGACGTAAGCCACGAAGACCCTGAGGTCAAGT
TCAACTGGTACGTGGACGGCGTGGAGGTGCATAATGCCAAGACAAA
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modified to include a heterologous signal sequence. (See, e.g., International Publication No. WO 96/34891.)

[0877] The amplified fragment is isolated from a 1% agarose gel using a commercially available kit ("GeneClean," BIO 101 Inc., La Jolla, Ca.). The fragment then is digested with appropriate restriction enzymes and again purified on a 1% agarose gel.

[0878] The amplified fragment is then digested with the same restriction enzyme and purified on a 1% agarose gel. The isolated fragment and the dephosphorylated vector are then ligated with T4 DNA ligase. *E. coli* HB101 or XL-1 Blue cells are then transformed and bacteria are identified that contain the fragment inserted into plasmid pC6 using, for instance, restriction enzyme analysis.

[0879] Chinese hamster ovary cells lacking an active DHFR gene is used for transfection. Five μ g of the expression plasmid pC6 or pC4 is cotransfected with 0.5 μ g of the plasmid pSVneo using lipofectin (Felgner et al., *supra*). The plasmid pSV2-neo contains a dominant selectable marker, the *neo* gene from Tn5 encoding an enzyme that confers resistance to a group of antibiotics including G418. The cells are seeded in alpha minus MEM supplemented with 1 mg/ml G418. After 2 days, the cells are trypsinized and seeded in hybridoma cloning plates (Greiner, Germany) in alpha minus MEM supplemented with 10, 25, or 50 ng/ml of methotrexate plus 1 mg/ml G418. After about 10-14 days single clones are trypsinized and then seeded in 6-well petri dishes or 10 ml flasks using different concentrations of methotrexate (50 nM, 100 nM, 200 nM, 400 nM, 800 nM). Clones growing at the highest concentrations of methotrexate are then transferred to new 6-well plates containing even higher concentrations of methotrexate (1 μ M, 2 μ M, 5 μ M, 10 mM, 20 mM). The same procedure is repeated until clones are obtained which grow at a concentration of 100 - 200 μ M. Expression of the desired gene product is analyzed, for instance, by SDS-PAGE and Western blot or by reversed phase HPLC analysis.

Example 9: Protein Fusions

[0880] The polypeptides of the present invention are preferably fused to other proteins. These fusion proteins can be used for a variety of applications. For example, fusion of the present polypeptides to His-tag, HA-tag, protein A, IgG domains, and maltose binding

[0872] Alternatively, the polypeptide can be expressed in stable cell lines containing the polynucleotide integrated into a chromosome. The co-transfection with a selectable marker such as DHFR, gpt, neomycin, or hygromycin allows the identification and isolation of the transfected cells.

[0873] The transfected gene can also be amplified to express large amounts of the encoded protein. The DHFR (dihydrofolate reductase) marker is useful in developing cell lines that carry several hundred or even several thousand copies of the gene of interest. (See, e.g., Alt, F. W., et al., J. Biol. Chem. 253:1357-1370 (1978); Hamlin, J. L. and Ma, C., Biochem. et Biophys. Acta, 1097:107-143 (1990); Page, M. J. and Sydenham, M. A., Biotechnology 9:64-68 (1991)). Another useful selection marker is the enzyme glutamine synthase (GS) (Murphy et al., Biochem J. 227:277-279 (1991); Bebbington et al., Bio/Technology 10:169-175 (1992)). Using these markers, the mammalian cells are grown in selective medium and the cells with the highest resistance are selected. These cell lines contain the amplified gene(s) integrated into a chromosome. Chinese hamster ovary (CHO) and NSO cells are often used for the production of proteins.

[0874] Derivatives of the plasmid pSV2-dhfr (ATCC Accession No. 37146), the expression vectors pC4 (ATCC Accession No. 209646) and pC6 (ATCC Accession No. 209647) contain the strong promoter (LTR) of the Rous Sarcoma Virus (Cullen et al., Molecular and Cellular Biology, 438-447 (March, 1985)) plus a fragment of the CMV-enhancer (Boshart et al., Cell 41:521-530 (1985)). Multiple cloning sites, e.g., with the restriction enzyme cleavage sites BamHI, XbaI and Asp718, facilitate the cloning of the gene of interest. The vectors also contain the 3' intron, the polyadenylation and termination signal of the rat preproinsulin gene, and the mouse DHFR gene under control of the SV40 early promoter.

[0875] Specifically, the plasmid pC6, for example, is digested with appropriate restriction enzymes and then dephosphorylated using calf intestinal phosphates by procedures known in the art. The vector is then isolated from a 1% agarose gel.

[0876] A polynucleotide of the present invention is amplified according to the protocol outlined in Example 1. If a naturally occurring signal sequence is used to produce the polypeptide of the present invention, the vector does not need a second signal peptide. Alternatively, if a naturally occurring signal sequence is not used, the vector can be

the recombinant baculovirus is used to infect Sf9 cells seeded in 35 mm dishes. Four days later the supernatants of these culture dishes are harvested and then they are stored at 4° C.

[0868] To verify the expression of the polypeptide, Sf9 cells are grown in Grace's medium supplemented with 10% heat-inactivated FBS. The cells are infected with the recombinant baculovirus containing the polynucleotide at a multiplicity of infection ("MOI") of about 2. If radiolabeled proteins are desired, 6 hours later the medium is removed and is replaced with SF900 II medium minus methionine and cysteine (available from Life Technologies Inc., Rockville, MD). After 42 hours, 5 μ Ci of 35 S-methionine and 5 μ Ci 35 S-cysteine (available from Amersham) are added. The cells are further incubated for 16 hours and then are harvested by centrifugation. The proteins in the supernatant as well as the intracellular proteins are analyzed by SDS-PAGE followed by autoradiography (if radiolabeled).

[0869] Microsequencing of the amino acid sequence of the amino terminus of purified protein may be used to determine the amino terminal sequence of the produced protein.

Example 8: Expression of a Polypeptide in Mammalian Cells

[0870] The polypeptide of the present invention can be expressed in a mammalian cell. A typical mammalian expression vector contains a promoter element, which mediates the initiation of transcription of mRNA, a protein coding sequence, and signals required for the termination of transcription and polyadenylation of the transcript. Additional elements include enhancers, Kozak sequences and intervening sequences flanked by donor and acceptor sites for RNA splicing. Highly efficient transcription is achieved with the early and late promoters from SV40, the long terminal repeats (LTRs) from Retroviruses, e.g., RSV, HTLV, HIV and the early promoter of the cytomegalovirus (CMV). However, cellular elements can also be used (e.g., the human actin promoter).

[0871] Suitable expression vectors for use in practicing the present invention include, for example, vectors such as pSVL and pMSG (Pharmacia, Uppsala, Sweden), pRSVcat (ATCC 37152), pSV2dhfr (ATCC 37146), pBC12MI (ATCC 67109), pCMVSPORT 2.0, and pCMVSPORT 3.0. Mammalian host cells that could be used include, human HeLa, 293, H9 and Jurkat cells, mouse NIH3T3 and C127 cells, Cos 1, Cos 7 and CV1, quail QC1-3 cells, mouse L cells and Chinese hamster ovary (CHO) cells.

[0864] The plasmid is digested with the corresponding restriction enzymes and optionally, can be dephosphorylated using calf intestinal phosphatase, using routine procedures known in the art. The DNA is then isolated from a 1% agarose gel using a commercially available kit ("GeneClean" BIO 101 Inc., La Jolla, Ca.).

[0865] The fragment and the dephosphorylated plasmid are ligated together with T4 DNA ligase. *E. coli* HB101 or other suitable *E. coli* hosts such as XL-1 Blue (Stratagene Cloning Systems, La Jolla, CA) cells are transformed with the ligation mixture and spread on culture plates. Bacteria containing the plasmid are identified by digesting DNA from individual colonies and analyzing the digestion product by gel electrophoresis. The sequence of the cloned fragment is confirmed by DNA sequencing.

[0866] Five μg of a plasmid containing the polynucleotide is co-transfected with 1.0 μg of a commercially available linearized baculovirus DNA ("BaculoGold™ baculovirus DNA, Pharmingen, San Diego, CA), using the lipofection method described by Felgner et al., Proc. Natl. Acad. Sci. USA 84:7413-7417 (1987). One μg of BaculoGold™ virus DNA and 5 μg of the plasmid are mixed in a sterile well of a microtiter plate containing 50 μl of serum-free Grace's medium (Life Technologies Inc., Gaithersburg, MD). Afterwards, 10 μl Lipofectin plus 90 μl Grace's medium are added, mixed and incubated for 15 minutes at room temperature. Then the transfection mixture is added drop-wise to Sf9 insect cells (ATCC CRL 1711) seeded in a 35 mm tissue culture plate with 1 ml Grace's medium without serum. The plate is then incubated for 5 hours at 27° C. The transfection solution is then removed from the plate and 1 ml of Grace's insect medium supplemented with 10% fetal calf serum is added. Cultivation is then continued at 27° C for four days.

[0867] After four days the supernatant is collected and a plaque assay is performed, as described by Summers and Smith, *supra*. An agarose gel with "Blue Gal" (Life Technologies Inc., Gaithersburg) is used to allow easy identification and isolation of gal-expressing clones, which produce blue-stained plaques. (A detailed description of a "plaque assay" of this type can also be found in the user's guide for insect cell culture and baculovirology distributed by Life Technologies Inc., Gaithersburg, page 9-10.) After appropriate incubation, blue stained plaques are picked with the tip of a micropipettor (e.g., Eppendorf). The agar containing the recombinant viruses is then resuspended in a microcentrifuge tube containing 200 μl of Grace's medium and the suspension containing

***Example 7: Cloning and Expression of a Polypeptide in a Baculovirus
Expression System***

[0860] In this example, the plasmid shuttle vector pA2 is used to insert a polynucleotide into a baculovirus to express a polypeptide. This expression vector contains the strong polyhedrin promoter of the *Autographa californica* nuclear polyhedrosis virus (AcMNPV) followed by convenient restriction sites such as BamHI, Xba I and Asp718. The polyadenylation site of the simian virus 40 ("SV40") is used for efficient polyadenylation. For easy selection of recombinant virus, the plasmid contains the beta-galactosidase gene from *E. coli* under control of a weak *Drosophila* promoter in the same orientation, followed by the polyadenylation signal of the polyhedrin gene. The inserted genes are flanked on both sides by viral sequences for cell-mediated homologous recombination with wild-type viral DNA to generate a viable virus that express the cloned polynucleotide.

[0861] Many other baculovirus vectors can be used in place of the vector above, such as pAc373, pVL941, and pAcIM1, as one skilled in the art would readily appreciate, as long as the construct provides appropriately located signals for transcription, translation, secretion and the like, including a signal peptide and an in-frame AUG as required. Such vectors are described, for instance, in Luckow et al., *Virology* 170:31-39 (1989).

[0862] Specifically, the cDNA sequence contained in the deposited clone, including the AUG initiation codon, is amplified using the PCR protocol described in Example 1. If a naturally occurring signal sequence is used to produce the polypeptide of the present invention, the pA2 vector does not need a second signal peptide. Alternatively, the vector can be modified (pA2 GP) to include a baculovirus leader sequence, using the standard methods described in Summers et al., "A Manual of Methods for Baculovirus Vectors and Insect Cell Culture Procedures," Texas Agricultural Experimental Station Bulletin No. 1555 (1987).

[0863] The amplified fragment is isolated from a 1% agarose gel using a commercially available kit ("GeneClean," BIO 101 Inc., La Jolla, Ca.). The fragment then is digested with appropriate restriction enzymes and again purified on a 1% agarose gel.

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[0855] The resulting washed inclusion bodies are solubilized with 1.5 M guanidine hydrochloride (GuHCl) for 2-4 hours. After 7000 xg centrifugation for 15 min., the pellet is discarded and the polypeptide containing supernatant is incubated at 4°C overnight to allow further GuHCl extraction.

[0856] Following high speed centrifugation (30,000 xg) to remove insoluble particles, the GuHCl solubilized protein is refolded by quickly mixing the GuHCl extract with 20 volumes of buffer containing 50 mM sodium, pH 4.5, 150 mM NaCl, 2 mM EDTA by vigorous stirring. The refolded diluted protein solution is kept at 4°C without mixing for 12 hours prior to further purification steps.

[0857] To clarify the refolded polypeptide solution, a previously prepared tangential filtration unit equipped with 0.16 µm membrane filter with appropriate surface area (e.g., Filtron), equilibrated with 40 mM sodium acetate, pH 6.0 is employed. The filtered sample is loaded onto a cation exchange resin (e.g., Poros HS-50, Perseptive Biosystems). The column is washed with 40 mM sodium acetate, pH 6.0 and eluted with 250 mM, 500 mM, 1000 mM, and 1500 mM NaCl in the same buffer, in a stepwise manner. The absorbance at 280 nm of the effluent is continuously monitored. Fractions are collected and further analyzed by SDS-PAGE.

[0858] Fractions containing the polypeptide are then pooled and mixed with 4 volumes of water. The diluted sample is then loaded onto a previously prepared set of tandem columns of strong anion (Poros HQ-50, Perseptive Biosystems) and weak anion (Poros CM-20, Perseptive Biosystems) exchange resins. The columns are equilibrated with 40 mM sodium acetate, pH 6.0. Both columns are washed with 40 mM sodium acetate, pH 6.0, 200 mM NaCl. The CM-20 column is then eluted using a 10 column volume linear gradient ranging from 0.2 M NaCl, 50 mM sodium acetate, pH 6.0 to 1.0 M NaCl, 50 mM sodium acetate, pH 6.5. Fractions are collected under constant A₂₈₀ monitoring of the effluent. Fractions containing the polypeptide (determined, for instance, by 16% SDS-PAGE) are then pooled.

[0859] The resultant polypeptide should exhibit greater than 95% purity after the above refolding and purification steps. No major contaminant bands should be observed from Commassie blue stained 16% SDS-PAGE gel when 5 µg of purified protein is loaded. The purified protein can also be tested for endotoxin/LPS contamination, and typically the LPS content is less than 0.1 ng/ml according to LAL assays.

neomycinphosphotransferase gene as a selection marker, 2) an *E. coli* origin of replication, 3) a T5 phage promoter sequence, 4) two lac operator sequences, 5) a Shine-Delgarno sequence, and 6) the lactose operon repressor gene (*lacIq*). The origin of replication (*oriC*) is derived from pUC19 (LTI, Gaithersburg, MD). The promoter and operator sequences are made synthetically.

[0850] DNA can be inserted into the pHE4a by restricting the vector with *NdeI* and *XbaI*, *BamHI*, *XhoI*, or *Asp718*, running the restricted product on a gel, and isolating the larger fragment (the stuffer fragment should be about 310 base pairs). The DNA insert is generated according to the PCR protocol described in Example 1, using PCR primers having restriction sites for *NdeI* (5' primer) and *XbaI*, *BamHI*, *XhoI*, or *Asp718* (3' primer). The PCR insert is gel purified and restricted with compatible enzymes. The insert and vector are ligated according to standard protocols.

[0851] The engineered vector could easily be substituted in the above protocol to express protein in a bacterial system.

Example 6: Purification of a Polypeptide from an Inclusion Body

[0852] The following alternative method can be used to purify a polypeptide expressed in *E. coli* when it is present in the form of inclusion bodies. Unless otherwise specified, all of the following steps are conducted at 4-10°C.

[0853] Upon completion of the production phase of the *E. coli* fermentation, the cell culture is cooled to 4-10°C and the cells harvested by continuous centrifugation at 15,000 rpm (Heraeus Sepatech). On the basis of the expected yield of protein per unit weight of cell paste and the amount of purified protein required, an appropriate amount of cell paste, by weight, is suspended in a buffer solution containing 100 mM Tris, 50 mM EDTA, pH 7.4. The cells are dispersed to a homogeneous suspension using a high shear mixer.

[0854] The cells are then lysed by passing the solution through a microfluidizer (Microfluidics, Corp. or APV Gaulin, Inc.) twice at 4000-6000 psi. The homogenate is then mixed with NaCl solution to a final concentration of 0.5 M NaCl, followed by centrifugation at 7000 xg for 15 min. The resultant pellet is washed again using 0.5M NaCl, 100 mM Tris, 50 mM EDTA, pH 7.4.

[0845] Clones containing the desired constructs are grown overnight (O/N) in liquid culture in LB media supplemented with both Amp (100 ug/ml) and Kan (25 ug/ml). The O/N culture is used to inoculate a large culture at a ratio of 1:100 to 1:250. The cells are grown to an optical density 600 (O.D.⁶⁰⁰) of between 0.4 and 0.6. IPTG (Isopropyl-B-D-thiogalacto pyranoside) is then added to a final concentration of 1 mM. IPTG induces by inactivating the lacI repressor, clearing the P/O leading to increased gene expression.

[0846] Cells are grown for an extra 3 to 4 hours. Cells are then harvested by centrifugation (20 mins at 6000Xg). The cell pellet is solubilized in the chaotropic agent 6 Molar Guanidine HCl by stirring for 3-4 hours at 4°C. The cell debris is removed by centrifugation, and the supernatant containing the polypeptide is loaded onto a nickel-nitrilo-tri-acetic acid ("Ni-NTA") affinity resin column (available from QIAGEN, Inc., *supra*). Proteins with a 6 x His tag bind to the Ni-NTA resin with high affinity and can be purified in a simple one-step procedure (for details see: The QIAexpressionist (1995) QIAGEN, Inc., *supra*).

[0847] Briefly, the supernatant is loaded onto the column in 6 M guanidine-HCl, pH 8. The column is first washed with 10 volumes of 6 M guanidine-HCl, pH 8, then washed with 10 volumes of 6 M guanidine-HCl pH 6, and finally the polypeptide is eluted with 6 M guanidine-HCl, pH 5.

[0848] The purified protein is then renatured by dialyzing it against phosphate-buffered saline (PBS) or 50 mM Na-acetate, pH 6 buffer plus 200 mM NaCl. Alternatively, the protein can be successfully refolded while immobilized on the Ni-NTA column. The recommended conditions are as follows: renature using a linear 6M-1M urea gradient in 500 mM NaCl, 20% glycerol, 20 mM Tris/HCl pH 7.4, containing protease inhibitors. The renaturation should be performed over a period of 1.5 hours or more. After renaturation the proteins are eluted by the addition of 250 mM imidazole. Imidazole is removed by a final dialyzing step against PBS or 50 mM sodium acetate pH 6 buffer plus 200 mM NaCl. The purified protein is stored at 4°C or frozen at -80°C.

[0849] In addition to the above expression vector, the present invention further includes an expression vector, called pHE4a (ATCC Accession Number 209645, deposited on February 25, 1998) which contains phage operator and promoter elements operatively linked to a polynucleotide of the present invention, called pHE4a. (ATCC Accession Number 209645, deposited on February 25, 1998.) This vector contains: 1) a

Example 4: Chromosomal Mapping of the Polynucleotides

[0842] An oligonucleotide primer set is designed according to the sequence at the 5' end of SEQ ID NO:X. This primer preferably spans about 100 nucleotides. This primer set is then used in a polymerase chain reaction under the following set of conditions: 30 seconds, 95°C; 1 minute, 56°C; 1 minute, 70°C. This cycle is repeated 32 times followed by one 5 minute cycle at 70°C. Human, mouse, and hamster DNA is used as template in addition to a somatic cell hybrid panel containing individual chromosomes or chromosome fragments (Bios, Inc). The reactions are analyzed on either 8% polyacrylamide gels or 3.5 % agarose gels. Chromosome mapping is determined by the presence of an approximately 100 bp PCR fragment in the particular somatic cell hybrid.

Example 5: Bacterial Expression of a Polypeptide

[0843] A polynucleotide encoding a polypeptide of the present invention is amplified using PCR oligonucleotide primers corresponding to the 5' and 3' ends of the DNA sequence, as outlined in Example 1, to synthesize insertion fragments. The primers used to amplify the cDNA insert should preferably contain restriction sites, such as BamHI and XbaI, at the 5' end of the primers in order to clone the amplified product into the expression vector. For example, BamHI and XbaI correspond to the restriction enzyme sites on the bacterial expression vector pQE-9. (Qiagen, Inc., Chatsworth, CA). This plasmid vector encodes antibiotic resistance (Amp^r), a bacterial origin of replication (ori), an IPTG-regulatable promoter/operator (P/O), a ribosome binding site (RBS), a 6-histidine tag (6-His), and restriction enzyme cloning sites.

[0844] The pQE-9 vector is digested with BamHI and XbaI and the amplified fragment is ligated into the pQE-9 vector maintaining the reading frame initiated at the bacterial RBS. The ligation mixture is then used to transform the E. coli strain M15/rep4 (Qiagen, Inc.) which contains multiple copies of the plasmid pREP4, which expresses the lacI repressor and also confers kanamycin resistance (Kan^r). Transformants are identified by their ability to grow on LB plates and ampicillin/kanamycin resistant colonies are selected. Plasmid DNA is isolated and confirmed by restriction analysis.

Example 3: Tissue specific expression analysis

[0838] The Human Genome Sciences, Inc. (HGS) database is derived from sequencing tissue and/or disease specific cDNA libraries. Libraries generated from a particular tissue are selected and the specific tissue expression pattern of EST groups or assembled contigs within these libraries is determined by comparison of the expression patterns of those groups or contigs within the entire database. ESTs and assembled contigs which show tissue specific expression are selected.

[0839] The original clone from which the specific EST sequence was generated, or in the case of an assembled contig, the clone from which the 5' most EST sequence was generated, is obtained from the catalogued library of clones and the insert amplified by PCR using methods known in the art. The PCR product is denatured and then transferred in 96 or 384 well format to a nylon membrane (Schleicher and Scheull) generating an array filter of tissue specific clones. Housekeeping genes, maize genes, and known tissue specific genes are included on the filters. These targets can be used in signal normalization and to validate assay sensitivity. Additional targets are included to monitor probe length and specificity of hybridization.

[0840] Radioactively labeled hybridization probes are generated by first strand cDNA synthesis per the manufacturer's instructions (Life Technologies) from mRNA/RNA samples prepared from the specific tissue being analyzed (e.g., prostate, prostate cancer, ovarian, ovarian cancer, etc.). The hybridization probes are purified by gel exclusion chromatography, quantitated, and hybridized with the array filters in hybridization bottles at 65°C overnight. The filters are washed under stringent conditions and signals are captured using a Fuji phosphorimager.

[0841] Data is extracted using AIS software and following background subtraction, signal normalization is performed. This includes a normalization of filter-wide expression levels between different experimental runs. Genes that are differentially expressed in the tissue of interest are identified.

art. For instance, a method similar to 5' RACE is available for generating the missing 5' end of a desired full-length transcript. (Fromont-Racine et al., Nucleic Acids Res. 21(7):1683-1684 (1993)).

[0834] Briefly, a specific RNA oligonucleotide is ligated to the 5' ends of a population of RNA presumably containing full-length gene RNA transcripts. A primer set containing a primer specific to the ligated RNA oligonucleotide and a primer specific to a known sequence of the gene of interest is used to PCR amplify the 5' portion of the desired full-length gene. This amplified product may then be sequenced and used to generate the full length gene.

[0835] This above method starts with total RNA isolated from the desired source, although poly-A+ RNA can be used. The RNA preparation can then be treated with phosphatase if necessary to eliminate 5' phosphate groups on degraded or damaged RNA which may interfere with the later RNA ligase step. The phosphatase should then be inactivated and the RNA treated with tobacco acid pyrophosphatase in order to remove the cap structure present at the 5' ends of messenger RNAs. This reaction leaves a 5' phosphate group at the 5' end of the cap cleaved RNA which can then be ligated to an RNA oligonucleotide using T4 RNA ligase.

[0836] This modified RNA preparation is used as a template for first strand cDNA synthesis using a gene specific oligonucleotide. The first strand synthesis reaction is used as a template for PCR amplification of the desired 5' end using a primer specific to the ligated RNA oligonucleotide and a primer specific to the known sequence of the gene of interest. The resultant product is then sequenced and analyzed to confirm that the 5' end sequence belongs to the desired gene.

Example 2: Isolation of Genomic Clones Corresponding to a Polynucleotide

[0837] A human genomic P1 library (Genomic Systems, Inc.) is screened by PCR using primers selected for the sequence corresponding to SEQ ID NO:X according to the method described in Example 1. (See also, Sambrook.)

2449. First, a plasmid is directly isolated by screening the clones using a polynucleotide probe corresponding to the nucleotide sequence of SEQ ID NO:X.

[0831] Particularly, a specific polynucleotide with 30-40 nucleotides is synthesized using an Applied Biosystems DNA synthesizer according to the sequence reported. The oligonucleotide is labeled, for instance, with ^{32}P - γ -ATP using T4 polynucleotide kinase and purified according to routine methods. (E.g., Maniatis et al., *Molecular Cloning: A Laboratory Manual*, Cold Spring Harbor Press, Cold Spring, NY (1982)). The plasmid mixture is transformed into a suitable host, as indicated above (such as XL-1 Blue (Stratagene)) using techniques known to those of skill in the art, such as those provided by the vector supplier or in related publications or patents cited above. The transformants are plated on 1.5% agar plates (containing the appropriate selection agent, e.g., ampicillin) to a density of about 150 transformants (colonies) per plate. These plates are screened using Nylon membranes according to routine methods for bacterial colony screening (e.g., Sambrook et al., *Molecular Cloning: A Laboratory Manual*, 2nd Edit., (1989), Cold Spring Harbor Laboratory Press, pages 1.93 to 1.104), or other techniques known to those of skill in the art.

[0832] Alternatively, two primers of 17-20 nucleotides derived from both ends of the nucleotide sequence of SEQ ID NO:X are synthesized and used to amplify the desired cDNA using the deposited cDNA plasmid as a template. The polymerase chain reaction is carried out under routine conditions, for instance, in 25 μl of reaction mixture with 0.5 μg of the above cDNA template. A convenient reaction mixture is 1.5-5 mM MgCl_2 , 0.01% (w/v) gelatin, 20 μM each of dATP, dCTP, dGTP, dTTP, 25 pmol of each primer and 0.25 Unit of Taq polymerase. Thirty five cycles of PCR (denaturation at 94°C for 1 min; annealing at 55°C for 1 min; elongation at 72°C for 1 min) are performed with a Perkin-Elmer Cetus automated thermal cycler. The amplified product is analyzed by agarose gel electrophoresis and the DNA band with expected molecular weight is excised and purified. The PCR product is verified to be the selected sequence by subcloning and sequencing the DNA product.

[0833] Several methods are available for the identification of the 5' or 3' non-coding portions of a gene which may not be present in the deposited clone. These methods include but are not limited to, filter probing, clone enrichment using specific probes, and protocols similar or identical to 5' and 3' "RACE" protocols which are well known in the

[0827] Vectors Lambda Zap (U.S. Patent Nos. 5,128,256 and 5,286,636), Uni-Zap XR (U.S. Patent Nos. 5,128, 256 and 5,286,636), Zap Express (U.S. Patent Nos. 5,128,256 and 5,286,636), pBluescript (pBS) (Short, J. M. et al., Nucleic Acids Res. 16:7583-7600 (1988); Altting-Mees, M. A. and Short, J. M., Nucleic Acids Res. 17:9494 (1989)) and pBK (Altting-Mees, M. A. et al., Strategies 5:58-61 (1992)) are commercially available from Stratagene Cloning Systems, Inc., 11011 N. Torrey Pines Road, La Jolla, CA, 92037. pBS contains an ampicillin resistance gene and pBK contains a neomycin resistance gene. Both can be transformed into E. coli strain XL-1 Blue, also available from Stratagene. pBS comes in 4 forms SK+, SK-, KS+ and KS. The S and K refers to the orientation of the polylinker to the T7 and T3 primer sequences which flank the polylinker region ("S" is for SacI and "K" is for KpnI which are the first sites on each respective end of the linker). "+" or "-" refer to the orientation of the f1 origin of replication ("ori"), such that in one orientation, single stranded rescue initiated from the f1 ori generates sense strand DNA and in the other, antisense.

[0828] Vectors pSport1, pCMVSPORT 2.0 and pCMVSPORT 3.0, were obtained from Life Technologies, Inc., P. O. Box 6009, Gaithersburg, MD 20897. All Sport vectors contain an ampicillin resistance gene and may be transformed into E. coli strain DH10B, also available from Life Technologies. (See, for instance, Gruber, C. E., et al., Focus 15:59 (1993)). Vector lafmid BA (Bento Soares, Columbia University, NY) contains an ampicillin resistance gene and can be transformed into E. coli strain XL-1 Blue. Vector pCR[®]2.1, which is available from Invitrogen, 1600 Faraday Avenue, Carlsbad, CA 92008, contains an ampicillin resistance gene and may be transformed into E. coli strain DH10B, available from Life Technologies. (See, for instance, Clark, J. M., Nuc. Acids Res. 16:9677-9686 (1988) and Mead, D. et al., Bio/Technology 9: (1991)). Preferably, a polynucleotide of the present invention does not comprise the phage vector sequences identified for the particular clone, as well as the corresponding plasmid vector sequences designated above.

[0829] The deposited material assigned the ATCC Deposit Number PTA-2448 and PTA-2449 contain at least a plasmid for each Clone ID NO:Z.

[0830] Two nonlimiting examples are provided below for isolating a particular clone from the deposited sample of plasmid cDNAs in ATCC Deposit No. PTA-2448 or PTA-

[0824] Also preferred is a method of treatment of an individual in need of a specific delivery of toxic compositions to diseased cells (e.g., tumors, leukemias or lymphomas), which method comprises administering to such an individual a Therapeutic comprising an amount of an isolated polypeptide of the invention, including, but not limited to a binding agent, or antibody of the claimed invention that are associated with toxin or cytotoxic prodrugs.

[0825] Having generally described the invention, the same will be more readily understood by reference to the following examples, which are provided by way of illustration and are not intended as limiting.

Examples

Example 1: Isolation of a Selected cDNA Clone From the Deposited Sample

[0826] Each Clone ID NO:Z is contained in a plasmid vector. In many cases, the vector used to construct the library is a phage vector from which a plasmid has been excised. The following correlates the related plasmid for each phage vector used in constructing the cDNA library. For example, where a particular clone is identified as being isolated in the vector "Lambda Zap," the corresponding deposited clone is in "pBluescript."

<u>Vector Used to Construct Library</u>	<u>Corresponding Deposited Plasmid</u>
Lambda Zap	pBluescript (pBS)
Uni-Zap XR	pBluescript (pBS)
Zap Express	pBK
lafmid BA	plafmid BA
pSport1	pSport1
pCMVSPORT 2.0	pCMVSPORT 2.0
pCMVSPORT 3.0	pCMVSPORT 3.0
pCR [®] 2.1	pCR [®] 2.1

[0819] Also preferred is a polypeptide molecule, wherein said polypeptide comprises an amino acid sequence selected from the group consisting of: polypeptide sequence of SEQ ID NO:Y; a polypeptide encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2; and a polypeptide encoded by the cDNA contained in Clone ID NO:Z.

[0820] Further preferred is a method of making a recombinant vector comprising inserting any of the above isolated nucleic acid molecule into a vector. Also preferred is the recombinant vector produced by this method. Also preferred is a method of making a recombinant host cell comprising introducing the vector into a host cell, as well as the recombinant host cell produced by this method.

[0821] Also preferred is a method of making an isolated polypeptide comprising culturing this recombinant host cell under conditions such that said polypeptide is expressed and recovering said polypeptide. Also preferred is this method of making an isolated polypeptide, wherein said recombinant host cell is a eukaryotic cell and said polypeptide is a human protein comprising an amino acid sequence selected from the group consisting of: polypeptide sequence of SEQ ID NO:Y; a polypeptide encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2; and a polypeptide encoded by the cDNA contained in Clone ID NO:Z. The isolated polypeptide produced by this method is also preferred.

[0822] Also preferred is a method of treatment of an individual in need of an increased level of a protein activity, which method comprises administering to such an individual a Therapeutic comprising an amount of an isolated polypeptide, polynucleotide, immunogenic fragment or analogue thereof, binding agent, antibody, or antigen binding fragment of the claimed invention effective to increase the level of said protein activity in said individual.

[0823] Also preferred is a method of treatment of an individual in need of a decreased level of a protein activity, which method comprised administering to such an individual a Therapeutic comprising an amount of an isolated polypeptide, polynucleotide, immunogenic fragment or analogue thereof, binding agent, antibody, or antigen binding fragment of the claimed invention effective to decrease the level of said protein activity in said individual.

sequence as defined in columns 8 and 9 of Table 2; and a polypeptide encoded by the cDNA contained in Clone ID NO:Z.

[0814] Also preferred is the above method for identifying the species, tissue or cell type of a biological sample, which method comprises a step of detecting polypeptide molecules comprising an amino acid sequence in a panel of at least two amino acid sequences, wherein at least one sequence in said panel is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the above group.

[0815] Also preferred is a method for diagnosing in a subject a pathological condition associated with abnormal structure or expression of a nucleic acid sequence identified in Table 1 or Table 2 encoding a polypeptide, which method comprises a step of detecting in a biological sample obtained from said subject polypeptide molecules comprising an amino acid sequence in a panel of at least two amino acid sequences, wherein at least one sequence in said panel is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: polypeptide sequence of SEQ ID NO:Y; a polypeptide encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2; and a polypeptide encoded by the cDNA contained in Clone ID NO:Z.

[0816] In any of these methods, the step of detecting said polypeptide molecules includes using an antibody.

[0817] Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a nucleotide sequence encoding a polypeptide wherein said polypeptide comprises an amino acid sequence that is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: polypeptide sequence of SEQ ID NO:Y; a polypeptide encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2; and a polypeptide encoded by the cDNA contained in Clone ID NO:Z.

[0818] Also preferred is an isolated nucleic acid molecule, wherein said nucleotide sequence encoding a polypeptide has been optimized for expression of said polypeptide in a prokaryotic host.

sequence as defined in columns 8 and 9 of Table 2; and a polypeptide encoded by the cDNA contained in Clone ID NO:Z.

[0810] Further preferred is a method for detecting in a biological sample a polypeptide comprising an amino acid sequence which is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: a polypeptide sequence of SEQ ID NO:Y; a polypeptide encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2; and a polypeptide encoded by the cDNA contained in Clone ID NO:Z; which method comprises a step of comparing an amino acid sequence of at least one polypeptide molecule in said sample with a sequence selected from said group and determining whether the sequence of said polypeptide molecule in said sample is at least 90% identical to said sequence of at least 10 contiguous amino acids.

[0811] Also preferred is the above method wherein said step of comparing an amino acid sequence of at least one polypeptide molecule in said sample with a sequence selected from said group comprises determining the extent of specific binding of polypeptides in said sample to an antibody which binds specifically to a polypeptide comprising an amino acid sequence that is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: a polypeptide sequence of SEQ ID NO:Y; a polypeptide encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2; and a polypeptide encoded by the cDNA contained in Clone ID NO:Z.

[0812] Also preferred is the above method wherein said step of comparing sequences is performed by comparing the amino acid sequence determined from a polypeptide molecule in said sample with said sequence selected from said group.

[0813] Also preferred is a method for identifying the species, tissue or cell type of a biological sample which method comprises a step of detecting polypeptide molecules in said sample, if any, comprising an amino acid sequence that is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: polypeptide sequence of SEQ ID NO:Y; a polypeptide encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide encoded by the nucleotide

amino acid sequence of SEQ ID NO:Y; a polypeptide encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2; and/or a polypeptide encoded by cDNA contained in Clone ID NO:Z.

[0803] Further preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to the complete amino acid sequence of SEQ ID NO:Y; a polypeptide encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2; and/or a polypeptide encoded by cDNA contained in Clone ID NO:Z.

[0804] Further preferred is an isolated polypeptide comprising an amino acid sequence at least 90% identical to a sequence of at least about 10 contiguous amino acids in the complete amino acid sequence of a polypeptide encoded by contained in Clone ID NO:Z

[0805] Also preferred is a polypeptide wherein said sequence of contiguous amino acids is included in the amino acid sequence of a portion of said polypeptide encoded by cDNA contained in Clone ID NO:Z; a polypeptide encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2; and/or the polypeptide sequence of SEQ ID NO:Y.

[0806] Also preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence of at least about 30 contiguous amino acids in the amino acid sequence of a polypeptide encoded by the cDNA contained in Clone ID NO:Z.

[0807] Also preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence of at least about 100 contiguous amino acids in the amino acid sequence of a polypeptide encoded by cDNA contained in Clone ID NO:Z.

[0808] Also preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to the amino acid sequence of a polypeptide encoded by the cDNA contained in Clone ID NO:Z.

[0809] Further preferred is an isolated antibody which binds specifically to a polypeptide comprising an amino acid sequence that is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: a polypeptide sequence of SEQ ID NO:Y; a polypeptide encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide encoded by the nucleotide

[0798] Also preferred is a composition of matter comprising isolated nucleic acid molecules wherein the nucleotide sequences of said nucleic acid molecules comprise a panel of at least two nucleotide sequences, wherein at least one sequence in said panel is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X or the complementary strand thereto; the nucleotide sequence as defined in column 5 of Table 1 or columns 8 and 9 of Table 2 or the complementary strand thereto; and a nucleotide sequence encoded by cDNA contained in Clone ID NO:Z. The nucleic acid molecules can comprise DNA molecules or RNA molecules.

[0799] Also preferred is a composition of matter comprising isolated nucleic acid molecules wherein the nucleotide sequences of said nucleic acid molecules comprise a DNA microarray or "chip" of at least 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 15, 20, 25, 30, 40, 50, 100, 150, 200, 250, 300, 500, 1000, 2000, 3000, or 4000 nucleotide sequences, wherein at least one sequence in said DNA microarray or "chip" is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1; and a nucleotide sequence encoded by a human cDNA clone identified by a cDNA "Clone ID" in Table 1.

[0800] Also preferred is an isolated polypeptide comprising an amino acid sequence at least 90% identical to a sequence of at least about 10 contiguous amino acids in the polypeptide sequence of SEQ ID NO:Y; a polypeptide encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2; and/or a polypeptide encoded by cDNA contained in Clone ID NO:Z.

[0801] Also preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence of at least about 30 contiguous amino acids in the amino acid sequence of SEQ ID NO:Y; a polypeptide encoded by SEQ ID NO:X or the complementary strand thereto; the polypeptide encoded by the nucleotide sequence as defined in columns 8 and 9 of Table 2; and/or a polypeptide encoded by cDNA contained in Clone ID NO:Z.

[0802] Further preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence of at least about 100 contiguous amino acids in the

from a nucleic acid molecule in said sample with said sequence selected from said group. The nucleic acid molecules can comprise DNA molecules or RNA molecules.

[0794] A further preferred embodiment is a method for identifying the species, tissue or cell type of a biological sample which method comprises a step of detecting nucleic acid molecules in said sample, if any, comprising a nucleotide sequence that is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X or the complementary strand thereto; the nucleotide sequence as defined in column 5 of Table 1 or columns 8 and 9 of Table 2 or the complementary strand thereto; and a nucleotide sequence of the cDNA contained in Clone ID NO:Z.

[0795] The method for identifying the species, tissue or cell type of a biological sample can comprise a step of detecting nucleic acid molecules comprising a nucleotide sequence in a panel of at least two nucleotide sequences, wherein at least one sequence in said panel is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from said group.

[0796] Also preferred is a method for diagnosing in a subject a pathological condition associated with abnormal structure or expression of a nucleotide sequence of SEQ ID NO:X or the complementary strand thereto; the nucleotide sequence as defined in column 5 of Table 1 or columns 8 and 9 of Table 2 or the complementary strand thereto; or the cDNA contained in Clone ID NO:Z which encodes a protein, wherein the method comprises a step of detecting in a biological sample obtained from said subject nucleic acid molecules, if any, comprising a nucleotide sequence that is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X or the complementary strand thereto; the nucleotide sequence as defined in column 5 of Table 1 or columns 8 and 9 of Table 2 or the complementary strand thereto; and a nucleotide sequence of cDNA contained in Clone ID NO:Z.

[0797] The method for diagnosing a pathological condition can comprise a step of detecting nucleic acid molecules comprising a nucleotide sequence in a panel of at least two nucleotide sequences, wherein at least one sequence in said panel is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from said group.

[0787] Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least 50 contiguous nucleotides of the cDNA sequence contained in Clone ID NO:Z.

[0788] Also preferred is an isolated nucleic acid molecule, wherein said sequence of at least 50 contiguous nucleotides is included in the nucleotide sequence of an open reading frame sequence encoded by cDNA contained in Clone ID NO:Z.

[0789] Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to sequence of at least 150 contiguous nucleotides in the nucleotide sequence encoded by cDNA contained in Clone ID NO:Z.

[0790] A further preferred embodiment is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to sequence of at least 500 contiguous nucleotides in the nucleotide sequence encoded by cDNA contained in Clone ID NO:Z.

[0791] A further preferred embodiment is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to the complete nucleotide sequence encoded by cDNA contained in Clone ID NO:Z.

[0792] A further preferred embodiment is a method for detecting in a biological sample a nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X or the complementary strand thereto; the nucleotide sequence as defined in column 5 of Table 1 or columns 8 and 9 of Table 2 or the complementary strand thereto; and a nucleotide sequence encoded by cDNA contained in Clone ID NO:Z; which method comprises a step of comparing a nucleotide sequence of at least one nucleic acid molecule in said sample with a sequence selected from said group and determining whether the sequence of said nucleic acid molecule in said sample is at least 95% identical to said selected sequence.

[0793] Also preferred is the above method wherein said step of comparing sequences comprises determining the extent of nucleic acid hybridization between nucleic acid molecules in said sample and a nucleic acid molecule comprising said sequence selected from said group. Similarly, also preferred is the above method wherein said step of comparing sequences is performed by comparing the nucleotide sequence determined

nucleotides in the nucleotide sequence of SEQ ID NO:X or the complementary strand thereto, the nucleotide sequence as defined in column 5 of Table 1 or columns 8 and 9 of Table 2 or the complementary strand thereto, and/or cDNA contained in Clone ID NO:Z.

[0781] Further preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least about 500 contiguous nucleotides in the nucleotide sequence of SEQ ID NO:X or the complementary strand thereto, the nucleotide sequence as defined in column 5 of Table 1 or columns 8 and 9 of Table 2 or the complementary strand thereto, and/or cDNA contained in Clone ID NO:Z.

[0782] A further preferred embodiment is a nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to the nucleotide sequence of the portion of SEQ ID NO:X defined in column 5, "ORF (From-To)", in Table 1.

[0783] A further preferred embodiment is a nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to the nucleotide sequence of the portion of SEQ ID NO:X defined in columns 8 and 9, "NT From" and "NT To", respectively, in Table 2.

[0784] A further preferred embodiment is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to the complete nucleotide sequence of SEQ ID NO:X or the complementary strand thereto, the nucleotide sequence as defined in column 5 of Table 1 or columns 8 and 9 of Table 2 or the complementary strand thereto, and/or cDNA contained in Clone ID NO:Z.

[0785] Also preferred is an isolated nucleic acid molecule which hybridizes under stringent hybridization conditions to a nucleic acid molecule comprising a nucleotide sequence of SEQ ID NO:X or the complementary strand thereto, the nucleotide sequence as defined in column 5 of Table 1 or columns 8 and 9 of Table 2 or the complementary strand thereto, and/or cDNA contained in Clone ID NO:Z, wherein said nucleic acid molecule which hybridizes does not hybridize under stringent hybridization conditions to a nucleic acid molecule having a nucleotide sequence consisting of only A residues or of only T residues.

[0786] Also preferred is a composition of matter comprising a DNA molecule which comprises the cDNA contained in Clone ID NO:Z.

[0774] A polypeptide, polynucleotide, agonist, or antagonist of the present invention may be used to change a mammal's mental state or physical state by influencing biorhythms, cardiac rhythms, depression (including depressive disorders), tendency for violence, tolerance for pain, reproductive capabilities (preferably by Activin or Inhibin-like activity), hormonal or endocrine levels, appetite, libido, memory, stress, or other cognitive qualities.

[0775] A polypeptide, polynucleotide, agonist, or antagonist of the present invention may also be used as a food additive or preservative, such as to increase or decrease storage capabilities, fat content, lipid, protein, carbohydrate, vitamins, minerals, cofactors or other nutritional components.

[0776] The above-recited applications have uses in a wide variety of hosts. Such hosts include, but are not limited to, human, murine, rabbit, goat, guinea pig, camel, horse, mouse, rat, hamster, pig, micro-pig, chicken, goat, cow, sheep, dog, cat, non-human primate, and human. In specific embodiments, the host is a mouse, rabbit, goat, guinea pig, chicken, rat, hamster, pig, sheep, dog or cat. In preferred embodiments, the host is a mammal. In most preferred embodiments, the host is a human.

Other Preferred Embodiments

[0777] Other preferred embodiments of the claimed invention include an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least about 50 contiguous nucleotides in the nucleotide sequence of SEQ ID NO:X or the complementary strand thereto, the nucleotide sequence as defined in column 5 of Table 1 or columns 8 and 9 of Table 2 or the complementary strand thereto, and/or cDNA contained in Clone ID NO:Z.

[0778] Also preferred is a nucleic acid molecule wherein said sequence of contiguous nucleotides is included in the nucleotide sequence of the portion of SEQ ID NO:X as defined in column 5, "ORF (From-To)", in Table 1.

[0779] Also preferred is a nucleic acid molecule wherein said sequence of contiguous nucleotides is included in the nucleotide sequence of the portion of SEQ ID NO:X as defined in columns 8 and 9, "NT From" and "NT To" respectively, in Table 2.

[0780] Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least about 150 contiguous

fibroblast cells and skeletal muscle cells, and therefore, facilitate the repair or replacement of damaged or diseased tissue.

[0768] A polypeptide, polynucleotide, agonist, or antagonist of the present invention may also be employed stimulate neuronal growth and to treat and prevent neuronal damage which occurs in certain neuronal disorders or neuro-degenerative conditions such as Alzheimer's disease, Parkinson's disease, and AIDS-related complex. A polypeptide, polynucleotide, agonist, or antagonist of the present invention may have the ability to stimulate chondrocyte growth, therefore, they may be employed to enhance bone and periodontal regeneration and aid in tissue transplants or bone grafts.

[0769] A polypeptide, polynucleotide, agonist, or antagonist of the present invention may be also be employed to prevent skin aging due to sunburn by stimulating keratinocyte growth.

[0770] A polypeptide, polynucleotide, agonist, or antagonist of the present invention may also be employed for preventing hair loss, since FGF family members activate hair-forming cells and promotes melanocyte growth. Along the same lines, a polypeptide, polynucleotide, agonist, or antagonist of the present invention may be employed to stimulate growth and differentiation of hematopoietic cells and bone marrow cells when used in combination with other cytokines.

[0771] A polypeptide, polynucleotide, agonist, or antagonist of the present invention may also be employed to maintain organs before transplantation or for supporting cell culture of primary tissues. A polypeptide, polynucleotide, agonist, or antagonist of the present invention may also be employed for inducing tissue of mesodermal origin to differentiate in early embryos.

[0772] A polypeptide, polynucleotide, agonist, or antagonist of the present invention may also increase or decrease the differentiation or proliferation of embryonic stem cells, besides, as discussed above, hematopoietic lineage.

[0773] A polypeptide, polynucleotide, agonist, or antagonist of the present invention may also be used to modulate mammalian characteristics, such as body height, weight, hair color, eye color, skin, percentage of adipose tissue, pigmentation, size, and shape (e.g., cosmetic surgery). Similarly, a polypeptide, polynucleotide, agonist, or antagonist of the present invention may be used to modulate mammalian metabolism affecting catabolism, anabolism, processing, utilization, and storage of energy.

combinatorial peptide libraries, or biased peptide libraries. The term "biased" is used herein to mean that the method of generating the library is manipulated so as to restrict one or more parameters that govern the diversity of the resulting collection of molecules, in this case peptides.

[0763] Thus, a truly random peptide library would generate a collection of peptides in which the probability of finding a particular amino acid at a given position of the peptide is the same for all 20 amino acids. A bias can be introduced into the library, however, by specifying, for example, that a lysine occur every fifth amino acid or that positions 4, 8, and 9 of a decapeptide library be fixed to include only arginine. Clearly, many types of biases can be contemplated, and the present invention is not restricted to any particular bias. Furthermore, the present invention contemplates specific types of peptide libraries, such as phage displayed peptide libraries and those that utilize a DNA construct comprising a lambda phage vector with a DNA insert.

[0764] As mentioned above, in the case of a binding molecule that is a polypeptide, the polypeptide may have about 6 to less than about 60 amino acid residues, preferably about 6 to about 10 amino acid residues, and most preferably, about 6 to about 22 amino acids. In another embodiment, a binding polypeptide has in the range of 15-100 amino acids, or 20-50 amino acids.

[0765] The selected binding polypeptide can be obtained by chemical synthesis or recombinant expression.

Other Activities

[0766] A polypeptide, polynucleotide, agonist, or antagonist of the present invention, as a result of the ability to stimulate vascular endothelial cell growth, may be employed in treatment for stimulating re-vascularization of ischemic tissues due to various disease conditions such as thrombosis, arteriosclerosis, and other cardiovascular conditions. The polypeptide, polynucleotide, agonist, or antagonist of the present invention may also be employed to stimulate angiogenesis and limb regeneration, as discussed above.

[0767] A polypeptide, polynucleotide, agonist, or antagonist of the present invention may also be employed for treating wounds due to injuries, burns, post-operative tissue repair, and ulcers since they are mitogenic to various cells of different origins, such as

might be the benzodiazepine structure.

[0758] Non-peptide oligomer libraries utilize a large number of monomers that are assembled together in ways that create new shapes that depend on the order of the monomers. Among the monomer units that have been used are carbamates, pyrrolinones, and morpholinos. Peptoids, peptide-like oligomers in which the side chain is attached to the alpha amino group rather than the alpha carbon, form the basis of another version of non-peptide oligomer libraries. The first non-peptide oligomer libraries utilized a single type of monomer and thus contained a repeating backbone. Recent libraries have utilized more than one monomer, giving the libraries added flexibility.

[0759] Screening the libraries can be accomplished by any of a variety of commonly known methods. See, e.g., the following references, which disclose screening of peptide libraries: Parmley and Smith, 1989, *Adv. Exp. Med. Biol.* 251:215-218; Scott and Smith, 1990, *Science* 249:386-390; Fowlkes et al., 1992, *BioTechniques* 13:422-427; Oldenburg et al., 1992, *Proc. Natl. Acad. Sci. USA* 89:5393-5397; Yu et al., 1994, *Cell* 76:933-945; Staudt et al., 1988, *Science* 241:577-580; Bock et al., 1992, *Nature* 355:564-566; Tuerk et al., 1992, *Proc. Natl. Acad. Sci. USA* 89:6988-6992; Ellington et al., 1992, *Nature* 355:850-852; U.S. Pat. No. 5,096,815, U.S. Pat. No. 5,223,409, and U.S. Pat. No. 5,198,346, all to Ladner et al.; Rebar and Pabo, 1993, *Science* 263:671-673; and CT Publication No. WO 94/18318.

[0760] In a specific embodiment, screening to identify a molecule that binds polypeptides of the invention can be carried out by contacting the library members with polypeptides of the invention immobilized on a solid phase and harvesting those library members that bind to the polypeptides of the invention. Examples of such screening methods, termed "panning" techniques are described by way of example in Parmley and Smith, 1988, *Gene* 73:305-318; Fowlkes et al., 1992, *BioTechniques* 13:422-427; PCT Publication No. WO 94/18318; and in references cited herein.

[0761] In another embodiment, the two-hybrid system for selecting interacting proteins in yeast (Fields and Song, 1989, *Nature* 340:245-246; Chien et al., 1991, *Proc. Natl. Acad. Sci. USA* 88:9578-9582) can be used to identify molecules that specifically bind to polypeptides of the invention.

[0762] Where the binding molecule is a polypeptide, the polypeptide can be conveniently selected from any peptide library, including random peptide libraries,

based libraries. Examples of chemically synthesized libraries are described in Fodor et al., 1991, *Science* 251:767-773; Houghten et al., 1991, *Nature* 354:84-86; Lam et al., 1991, *Nature* 354:82-84; Medynski, 1994, *Bio/Technology* 12:709-710; Gallop et al., 1994, *J. Medicinal Chemistry* 37(9):1233-1251; Ohlmeyer et al., 1993, *Proc. Natl. Acad. Sci. USA* 90:10922-10926; Erb et al., 1994, *Proc. Natl. Acad. Sci. USA* 91:11422-11426; Houghten et al., 1992, *Biotechniques* 13:412; Jayawickreme et al., 1994, *Proc. Natl. Acad. Sci. USA* 91:1614-1618; Salmon et al., 1993, *Proc. Natl. Acad. Sci. USA* 90:11708-11712; PCT Publication No. WO 93/20242; and Brenner and Lerner, 1992, *Proc. Natl. Acad. Sci. USA* 89:5381-5383.

[0753] Examples of phage display libraries are described in Scott and Smith, 1990, *Science* 249:386-390; Devlin et al., 1990, *Science*, 249:404-406; Christian, R. B., et al., 1992, *J. Mol. Biol.* 227:711-718; Lenstra, 1992, *J. Immunol. Meth.* 152:149-157; Kay et al., 1993, *Gene* 128:59-65; and PCT Publication No. WO 94/18318 dated Aug. 18, 1994.

[0754] In vitro translation-based libraries include but are not limited to those described in PCT Publication No. WO 91/05058 dated Apr. 18, 1991; and Mattheakis et al., 1994, *Proc. Natl. Acad. Sci. USA* 91:9022-9026.

[0755] By way of examples of nonpeptide libraries, a benzodiazepine library (see e.g., Bunin et al., 1994, *Proc. Natl. Acad. Sci. USA* 91:4708-4712) can be adapted for use. Peptoid libraries (Simon et al., 1992, *Proc. Natl. Acad. Sci. USA* 89:9367-9371) can also be used. Another example of a library that can be used, in which the amide functionalities in peptides have been permethylated to generate a chemically transformed combinatorial library, is described by Ostresh et al. (1994, *Proc. Natl. Acad. Sci. USA* 91:11138-11142).

[0756] The variety of non-peptide libraries that are useful in the present invention is great. For example, Ecker and Crooke, 1995, *Bio/Technology* 13:351-360 list benzodiazepines, hydantoins, piperazinediones, biphenyls, sugar analogs, beta-mercaptoketones, arylacetic acids, acylpiperidines, benzopyrans, cubanes, xanthines, aminimides, and oxazolones as among the chemical species that form the basis of various libraries.

[0757] Non-peptide libraries can be classified broadly into two types: decorated monomers and oligomers. Decorated monomer libraries employ a relatively simple scaffold structure upon which a variety functional groups is added. Often the scaffold will be a molecule with a known useful pharmacological activity. For example, the scaffold

[0750] Alternatively, one may also separate a plurality of polypeptides into substantially separate fractions comprising a subset of or individual polypeptides. For instance, one can separate the plurality of polypeptides by gel electrophoresis, column chromatography, or like method known to those of ordinary skill for the separation of polypeptides. The individual polypeptides can also be produced by a transformed host cell in such a way as to be expressed on or about its outer surface (e.g., a recombinant phage). Individual isolates can then be "probed" by the polypeptides of the invention, optionally in the presence of an inducer should one be required for expression, to determine if any selective affinity interaction takes place between the polypeptides and the individual clone. Prior to contacting the polypeptides with each fraction comprising individual polypeptides, the polypeptides could first be transferred to a solid support for additional convenience. Such a solid support may simply be a piece of filter membrane, such as one made of nitrocellulose or nylon. In this manner, positive clones could be identified from a collection of transformed host cells of an expression library, which harbor a DNA construct encoding a polypeptide having a selective affinity for polypeptides of the invention. Furthermore, the amino acid sequence of the polypeptide having a selective affinity for the polypeptides of the invention can be determined directly by conventional means or the coding sequence of the DNA encoding the polypeptide can frequently be determined more conveniently. The primary sequence can then be deduced from the corresponding DNA sequence. If the amino acid sequence is to be determined from the polypeptide itself, one may use microsequencing techniques. The sequencing technique may include mass spectroscopy.

[0751] In certain situations, it may be desirable to wash away any unbound polypeptides from a mixture of the polypeptides of the invention and the plurality of polypeptides prior to attempting to determine or to detect the presence of a selective affinity interaction. Such a wash step may be particularly desirable when the polypeptides of the invention or the plurality of polypeptides are bound to a solid support.

[0752] The plurality of molecules provided according to this method may be provided by way of diversity libraries, such as random or combinatorial peptide or nonpeptide libraries which can be screened for molecules that specifically bind polypeptides of the invention. Many libraries are known in the art that can be used, e.g., chemically synthesized libraries, recombinant (e.g., phage display libraries), and in vitro translation-

[0744] The antagonist/agonist may also be employed to prevent the growth of scar tissue during wound healing.

[0745] The antagonist/agonist may also be employed to treat the diseases described herein.

[0746] Thus, the invention provides a method of treating disorders or diseases, including but not limited to the disorders or diseases listed throughout this application, associated with overexpression of a polynucleotide of the present invention by administering to a patient (a) an antisense molecule directed to the polynucleotide of the present invention, and/or (b) a ribozyme directed to the polynucleotide of the present invention.

Binding Peptides and Other Molecules

[0747] The invention also encompasses screening methods for identifying polypeptides and nonpolypeptides that bind polypeptides of the invention, and the binding molecules identified thereby. These binding molecules are useful, for example, as agonists and antagonists of the polypeptides of the invention. Such agonists and antagonists can be used, in accordance with the invention, in the therapeutic embodiments described in detail, below.

[0748] This method comprises the steps of:

contacting polypeptides of the invention with a plurality of molecules; and

identifying a molecule that binds the polypeptides of the invention.

[0749] The step of contacting the polypeptides of the invention with the plurality of molecules may be effected in a number of ways. For example, one may contemplate immobilizing the polypeptides on a solid support and bringing a solution of the plurality of molecules in contact with the immobilized polypeptides. Such a procedure would be akin to an affinity chromatographic process, with the affinity matrix being comprised of the immobilized polypeptides of the invention. The molecules having a selective affinity for the polypeptides can then be purified by affinity selection. The nature of the solid support, process for attachment of the polypeptides to the solid support, solvent, and conditions of the affinity isolation or selection are largely conventional and well known to those of ordinary skill in the art.

[0740] Potential antagonists according to the invention also include catalytic RNA, or a ribozyme (See, e.g., PCT International Publication WO 90/11364, published October 4, 1990; Sarver et al, Science 247:1222-1225 (1990). While ribozymes that cleave mRNA at site specific recognition sequences can be used to destroy mRNAs, the use of hammerhead ribozymes is preferred. Hammerhead ribozymes cleave mRNAs at locations dictated by flanking regions that form complementary base pairs with the target mRNA. The sole requirement is that the target mRNA have the following sequence of two bases: 5'-UG-3'. The construction and production of hammerhead ribozymes is well known in the art and is described more fully in Haseloff and Gerlach, Nature 334:585-591 (1988). There are numerous potential hammerhead ribozyme cleavage sites within the nucleotide sequence of SEQ ID NO:X. Preferably, the ribozyme is engineered so that the cleavage recognition site is located near the 5' end of the mRNA; i.e., to increase efficiency and minimize the intracellular accumulation of non-functional mRNA transcripts.

[0741] As in the antisense approach, the ribozymes of the invention can be composed of modified oligonucleotides (e.g., for improved stability, targeting, etc.) and should be delivered to cells which express *in vivo*. DNA constructs encoding the ribozyme may be introduced into the cell in the same manner as described above for the introduction of antisense encoding DNA. A preferred method of delivery involves using a DNA construct "encoding" the ribozyme under the control of a strong constitutive promoter, such as, for example, pol III or pol II promoter, so that transfected cells will produce sufficient quantities of the ribozyme to destroy endogenous messages and inhibit translation. Since ribozymes unlike antisense molecules, are catalytic, a lower intracellular concentration is required for efficiency.

[0742] Antagonist/agonist compounds may be employed to inhibit the cell growth and proliferation effects of the polypeptides of the present invention on neoplastic cells and tissues, i.e. stimulation of angiogenesis of tumors, and, therefore, retard or prevent abnormal cellular growth and proliferation, for example, in tumor formation or growth.

[0743] The antagonist/agonist may also be employed to prevent hyper-vascular diseases, and prevent the proliferation of epithelial lens cells after extracapsular cataract surgery. Prevention of the mitogenic activity of the polypeptides of the present invention may also be desirous in cases such as restenosis after balloon angioplasty.

2-methyladenine, 2-methylguanine, 3-methylcytosine, 5-methylcytosine, N6-adenine, 7-methylguanine, 5-methylaminomethyluracil, 5-methoxyaminomethyl-2-thiouracil, beta-D-mannosylqueosine, 5'-methoxycarboxymethyluracil, 5-methoxyuracil, 2-methylthio-N6-isopentenyladenine, uracil-5-oxyacetic acid (v), wybutoxosine, pseudouracil, queosine, 2-thiocytosine, 5-methyl-2-thiouracil, 2-thiouracil, 4-thiouracil, 5-methyluracil, uracil-5-oxyacetic acid methylester, uracil-5-oxyacetic acid (v), 5-methyl-2-thiouracil, 3-(3-amino-3-N-2-carboxypropyl) uracil, (acp3)w, and 2,6-diaminopurine.

[0735] The antisense oligonucleotide may also comprise at least one modified sugar moiety selected from the group including, but not limited to, arabinose, 2-fluoroarabinose, xylulose, and hexose.

[0736] In yet another embodiment, the antisense oligonucleotide comprises at least one modified phosphate backbone selected from the group including, but not limited to, a phosphorothioate, a phosphorodithioate, a phosphoramidothioate, a phosphoramidate, a phosphordiamidate, a methylphosphonate, an alkyl phosphotriester, and a formacetal or analog thereof.

[0737] In yet another embodiment, the antisense oligonucleotide is an a-anomeric oligonucleotide. An a-anomeric oligonucleotide forms specific double-stranded hybrids with complementary RNA in which, contrary to the usual b-units, the strands run parallel to each other (Gautier et al., 1987, Nucl. Acids Res. 15:6625-6641). The oligonucleotide is a 2'-O-methylribonucleotide (Inoue et al., 1987, Nucl. Acids Res. 15:6131-6148), or a chimeric RNA-DNA analogue (Inoue et al., 1987, FEBS Lett. 215:327-330).

[0738] Polynucleotides of the invention may be synthesized by standard methods known in the art, e.g. by use of an automated DNA synthesizer (such as are commercially available from Biosearch, Applied Biosystems, etc.). As examples, phosphorothioate oligonucleotides may be synthesized by the method of Stein et al. (1988, Nucl. Acids Res. 16:3209), methylphosphonate oligonucleotides can be prepared by use of controlled pore glass polymer supports (Sarin et al., 1988, Proc. Natl. Acad. Sci. U.S.A. 85:7448-7451), etc.

[0739] While antisense nucleotides complementary to the coding region sequence could be used, those complementary to the transcribed untranslated region are most preferred.

translation of mRNAs as well. See generally, Wagner, R., 1994, Nature 372:333-335. Thus, oligonucleotides complementary to either the 5'- or 3'- non- translated, non-coding regions of polynucleotide sequences described herein could be used in an antisense approach to inhibit translation of endogenous mRNA. Oligonucleotides complementary to the 5' untranslated region of the mRNA should include the complement of the AUG start codon. Antisense oligonucleotides complementary to mRNA coding regions are less efficient inhibitors of translation but could be used in accordance with the invention. Whether designed to hybridize to the 5'-, 3'- or coding region of mRNA of the present invention, antisense nucleic acids should be at least six nucleotides in length, and are preferably oligonucleotides ranging from 6 to about 50 nucleotides in length. In specific aspects the oligonucleotide is at least 10 nucleotides, at least 17 nucleotides, at least 25 nucleotides or at least 50 nucleotides.

[0733] The polynucleotides of the invention can be DNA or RNA or chimeric mixtures or derivatives or modified versions thereof, single-stranded or double-stranded. The oligonucleotide can be modified at the base moiety, sugar moiety, or phosphate backbone, for example, to improve stability of the molecule, hybridization, etc. The oligonucleotide may include other appended groups such as peptides (e.g., for targeting host cell receptors *in vivo*), or agents facilitating transport across the cell membrane (see, e.g., Letsinger et al., 1989, Proc. Natl. Acad. Sci. U.S.A. 86:6553-6556; Lemaitre et al., 1987, Proc. Natl. Acad. Sci. 84:648-652; PCT Publication No. WO88/09810, published December 15, 1988) or the blood-brain barrier (see, e.g., PCT Publication No. WO89/10134, published April 25, 1988), hybridization-triggered cleavage agents. (See, e.g., Krol et al., 1988, BioTechniques 6:958-976) or intercalating agents. (See, e.g., Zon, 1988, Pharm. Res. 5:539-549). To this end, the oligonucleotide may be conjugated to another molecule, e.g., a peptide, hybridization triggered cross-linking agent, transport agent, hybridization-triggered cleavage agent, etc.

[0734] The antisense oligonucleotide may comprise at least one modified base moiety which is selected from the group including, but not limited to, 5-fluorouracil, 5-bromouracil, 5-chlorouracil, 5-iodouracil, hypoxanthine, xantine, 4-acetylcytosine, 5-(carboxyhydroxymethyl) uracil, 5-carboxymethylaminomethyl-2-thiouridine, 5-carboxymethylaminomethyluracil, dihydrouracil, beta-D-galactosylqueosine, inosine, N6-isopentenyladenine, 1-methylguanine, 1-methylinosine, 2,2-dimethylguanine,

[0730] In one embodiment, the antisense nucleic acid of the invention is produced intracellularly by transcription from an exogenous sequence. For example, a vector or a portion thereof, is transcribed, producing an antisense nucleic acid (RNA) of the invention. Such a vector would contain a sequence encoding the antisense nucleic acid. Such a vector can remain episomal or become chromosomally integrated, as long as it can be transcribed to produce the desired antisense RNA. Such vectors can be constructed by recombinant DNA technology methods standard in the art. Vectors can be plasmid, viral, or others known in the art, used for replication and expression in vertebrate cells. Expression of the sequence encoding the polypeptide of the present invention or fragments thereof, can be by any promoter known in the art to act in vertebrate, preferably human cells. Such promoters can be inducible or constitutive. Such promoters include, but are not limited to, the SV40 early promoter region (Bernoist and Chambon, *Nature* 29:304-310 (1981), the promoter contained in the 3' long terminal repeat of Rous sarcoma virus (Yamamoto et al., *Cell* 22:787-797 (1980), the herpes thymidine promoter (Wagner et al., *Proc. Natl. Acad. Sci. U.S.A.* 78:1441-1445 (1981), the regulatory sequences of the metallothionein gene (Brinster, et al., *Nature* 296:39-42 (1982)), etc.

[0731] The antisense nucleic acids of the invention comprise a sequence complementary to at least a portion of an RNA transcript of a gene of the present invention. However, absolute complementarity, although preferred, is not required. A sequence "complementary to at least a portion of an RNA," referred to herein, means a sequence having sufficient complementarity to be able to hybridize with the RNA, forming a stable duplex; in the case of double stranded antisense nucleic acids, a single strand of the duplex DNA may thus be tested, or triplex formation may be assayed. The ability to hybridize will depend on both the degree of complementarity and the length of the antisense nucleic acid. Generally, the larger the hybridizing nucleic acid, the more base mismatches with a RNA it may contain and still form a stable duplex (or triplex as the case may be). One skilled in the art can ascertain a tolerable degree of mismatch by use of standard procedures to determine the melting point of the hybridized complex.

[0732] Oligonucleotides that are complementary to the 5' end of the message, e.g., the 5' untranslated sequence up to and including the AUG initiation codon, should work most efficiently at inhibiting translation. However, sequences complementary to the 3' untranslated sequences of mRNAs have been shown to be effective at inhibiting

complementary strand thereof, and/or to cDNA sequences contained in cDNA Clone ID NO:Z identified for example, in Table 1. In one embodiment, antisense sequence is generated internally, by the organism, in another embodiment, the antisense sequence is separately administered (see, for example, O'Connor, J., *Neurochem.* 56:560 (1991). *Oligodeoxynucleotides as Antisense Inhibitors of Gene Expression*, CRC Press, Boca Raton, FL (1988). Antisense technology can be used to control gene expression through antisense DNA or RNA, or through triple-helix formation. Antisense techniques are discussed for example, in Okano, J., *Neurochem.* 56:560 (1991); *Oligodeoxynucleotides as Antisense Inhibitors of Gene Expression*, CRC Press, Boca Raton, FL (1988). Triple helix formation is discussed in, for instance, Lee et al., *Nucleic Acids Research* 6:3073 (1979); Cooney et al., *Science* 241:456 (1988); and Dervan et al., *Science* 251:1300 (1991). The methods are based on binding of a polynucleotide to a complementary DNA or RNA.

[0728] For example, the use of c-myc and c-myb antisense RNA constructs to inhibit the growth of the non-lymphocytic leukemia cell line HL-60 and other cell lines was previously described. (Wickstrom et al. (1988); Anfossi et al. (1989)). These experiments were performed *in vitro* by incubating cells with the oligoribonucleotide. A similar procedure for *in vivo* use is described in WO 91/15580. Briefly, a pair of oligonucleotides for a given antisense RNA is produced as follows: A sequence complimentary to the first 15 bases of the open reading frame is flanked by an EcoR1 site on the 5' end and a HindIII site on the 3' end. Next, the pair of oligonucleotides is heated at 90°C for one minute and then annealed in 2X ligation buffer (20mM TRIS HCl pH 7.5, 10mM MgCl₂, 10mM dithiothreitol (DTT) and 0.2 mM ATP) and then ligated to the EcoR1/Hind III site of the retroviral vector PMV7 (WO 91/15580).

[0729] For example, the 5' coding portion of a polynucleotide that encodes the polypeptide of the present invention may be used to design an antisense RNA oligonucleotide of from about 10 to 40 base pairs in length. A DNA oligonucleotide is designed to be complementary to a region of the gene involved in transcription thereby preventing transcription and the production of the receptor. The antisense RNA oligonucleotide hybridizes to the mRNA *in vivo* and blocks translation of the mRNA molecule into receptor polypeptide.

which are stably transformed with recombinant nucleic acids expressing the polypeptide or fragment. Drugs are screened against such transformed cells in competitive binding assays. One may measure, for example, the formulation of complexes between the agent being tested and a polypeptide of the present invention.

[0724] Thus, the present invention provides methods of screening for drugs or any other agents which affect activities mediated by the polypeptides of the present invention. These methods comprise contacting such an agent with a polypeptide of the present invention or a fragment thereof and assaying for the presence of a complex between the agent and the polypeptide or a fragment thereof, by methods well known in the art. In such a competitive binding assay, the agents to screen are typically labeled. Following incubation, free agent is separated from that present in bound form, and the amount of free or uncomplexed label is a measure of the ability of a particular agent to bind to the polypeptides of the present invention.

[0725] Another technique for drug screening provides high throughput screening for compounds having suitable binding affinity to the polypeptides of the present invention, and is described in great detail in European Patent Application 84/03564, published on September 13, 1984, which is incorporated herein by reference herein. Briefly stated, large numbers of different small peptide test compounds are synthesized on a solid substrate, such as plastic pins or some other surface. The peptide test compounds are reacted with polypeptides of the present invention and washed. Bound polypeptides are then detected by methods well known in the art. Purified polypeptides are coated directly onto plates for use in the aforementioned drug screening techniques. In addition, non-neutralizing antibodies may be used to capture the peptide and immobilize it on the solid support.

[0726] This invention also contemplates the use of competitive drug screening assays in which neutralizing antibodies capable of binding polypeptides of the present invention specifically compete with a test compound for binding to the polypeptides or fragments thereof. In this manner, the antibodies are used to detect the presence of any peptide which shares one or more antigenic epitopes with a polypeptide of the invention.

Antisense And Ribozyme (Antagonists)

[0727] In specific embodiments, antagonists according to the present invention are nucleic acids corresponding to the sequences contained in SEQ ID NO:X, or the

[0720] In another embodiment, the invention provides a method for the specific destruction of cells (e.g., the destruction of tumor cells) by administering polypeptides of the invention (e.g., polypeptides of the invention or antibodies of the invention) in association with toxins or cytotoxic prodrugs.

[0721] By "toxin" is meant compounds that bind and activate endogenous cytotoxic effector systems, radioisotopes, holotoxins, modified toxins, catalytic subunits of toxins, or any molecules or enzymes not normally present in or on the surface of a cell that under defined conditions cause the cell's death. Toxins that may be used according to the methods of the invention include, but are not limited to, radioisotopes known in the art, compounds such as, for example, antibodies (or complement fixing containing portions thereof) that bind an inherent or induced endogenous cytotoxic effector system, thymidine kinase, endonuclease, RNase, alpha toxin, ricin, abrin, *Pseudomonas* exotoxin A, diphtheria toxin, saporin, momordin, gelonin, pokeweed antiviral protein, alpha-sarcin and cholera toxin. By "cytotoxic prodrug" is meant a non-toxic compound that is converted by an enzyme, normally present in the cell, into a cytotoxic compound. Cytotoxic prodrugs that may be used according to the methods of the invention include, but are not limited to, glutamyl derivatives of benzoic acid mustard alkylating agent, phosphate derivatives of etoposide or mitomycin C, cytosine arabinoside, daunorubisin, and phenoxacetamide derivatives of doxorubicin.

Drug Screening

[0722] Further contemplated is the use of the polypeptides of the present invention, or the polynucleotides encoding these polypeptides, to screen for molecules which modify the activities of the polypeptides of the present invention. Such a method would include contacting the polypeptide of the present invention with a selected compound(s) suspected of having antagonist or agonist activity, and assaying the activity of these polypeptides following binding.

[0723] This invention is particularly useful for screening therapeutic compounds by using the polypeptides of the present invention, or binding fragments thereof, in any of a variety of drug screening techniques. The polypeptide or fragment employed in such a test may be affixed to a solid support, expressed on a cell surface, free in solution, or located intracellularly. One method of drug screening utilizes eukaryotic or prokaryotic host cells

a second messenger response is measured to determine if the compound is a potential agonist or antagonist. Such second messenger systems include but are not limited to, cAMP guanylate cyclase, ion channels or phosphoinositide hydrolysis.

[0716] All of these above assays can be used as diagnostic or prognostic markers. The molecules discovered using these assays can be used to treat disease or to bring about a particular result in a patient (e.g., blood vessel growth) by activating or inhibiting the polypeptide/molecule. Moreover, the assays can discover agents which may inhibit or enhance the production of the polypeptides of the invention from suitably manipulated cells or tissues.

[0717] Therefore, the invention includes a method of identifying compounds which bind to a polypeptide of the invention comprising the steps of: (a) incubating a candidate binding compound with a polypeptide of the present invention; and (b) determining if binding has occurred. Moreover, the invention includes a method of identifying agonists/antagonists comprising the steps of: (a) incubating a candidate compound with a polypeptide of the present invention, (b) assaying a biological activity, and (b) determining if a biological activity of the polypeptide has been altered.

Targeted Delivery

[0718] In another embodiment, the invention provides a method of delivering compositions to targeted cells expressing a receptor for a polypeptide of the invention, or cells expressing a cell bound form of a polypeptide of the invention.

[0719] As discussed herein, polypeptides or antibodies of the invention may be associated with heterologous polypeptides, heterologous nucleic acids, toxins, or prodrugs via hydrophobic, hydrophilic, ionic and/or covalent interactions. In one embodiment, the invention provides a method for the specific delivery of compositions of the invention to cells by administering polypeptides of the invention (including antibodies) that are associated with heterologous polypeptides or nucleic acids. In one example, the invention provides a method for delivering a therapeutic protein into the targeted cell. In another example, the invention provides a method for delivering a single stranded nucleic acid (e.g., antisense or ribozymes) or double stranded nucleic acid (e.g., DNA that can integrate into the cell's genome or replicate episomally and that can be transcribed) into the targeted cell.

be recombined with one or more components, motifs, sections, parts, domains, fragments, etc. of one or more heterologous molecules. In preferred embodiments, the heterologous molecules are family members. In further preferred embodiments, the heterologous molecule is a growth factor such as, for example, platelet-derived growth factor (PDGF), insulin-like growth factor (IGF-I), transforming growth factor (TGF)-alpha, epidermal growth factor (EGF), fibroblast growth factor (FGF), TGF-beta, bone morphogenetic protein (BMP)-2, BMP-4, BMP-5, BMP-6, BMP-7, activins A and B, decapentaplegic(dpp), 60A, OP-2, dorsalin, growth differentiation factors (GDFs), nodal, MIS, inhibin-alpha, TGF-beta1, TGF-beta2, TGF-beta3, TGF-beta5, and glial-derived neurotrophic factor (GDNF).

[0713] Other preferred fragments are biologically active fragments of the polypeptide of the present invention. Biologically active fragments are those exhibiting activity similar, but not necessarily identical, to an activity of the polypeptide of the present invention. The biological activity of the fragments may include an improved desired activity, or a decreased undesirable activity.

[0714] Additionally, this invention provides a method of screening compounds to identify those which modulate the action of the polypeptide of the present invention. An example of such an assay comprises combining a mammalian fibroblast cell, a the polypeptide of the present invention, the compound to be screened and $^3\text{[H]}$ thymidine under cell culture conditions where the fibroblast cell would normally proliferate. A control assay may be performed in the absence of the compound to be screened and compared to the amount of fibroblast proliferation in the presence of the compound to determine if the compound stimulates proliferation by determining the uptake of $^3\text{[H]}$ thymidine in each case. The amount of fibroblast cell proliferation is measured by liquid scintillation chromatography which measures the incorporation of $^3\text{[H]}$ thymidine. Both agonist and antagonist compounds may be identified by this procedure.

[0715] In another method, a mammalian cell or membrane preparation expressing a receptor for a polypeptide of the present invention is incubated with a labeled polypeptide of the present invention in the presence of the compound. The ability of the compound to enhance or block this interaction could then be measured. Alternatively, the response of a known second messenger system following interaction of a compound to be screened and the receptor is measured and the ability of the compound to bind to the receptor and elicit

Transfected cells which are grown on glass slides are exposed to the polypeptide of the present invention, after they have been labeled. The polypeptides can be labeled by a variety of means including iodination or inclusion of a recognition site for a site-specific protein kinase.

[0710] Following fixation and incubation, the slides are subjected to auto-radiographic analysis. Positive pools are identified and sub-pools are prepared and re-transfected using an iterative sub-pooling and re-screening process, eventually yielding a single clones that encodes the putative receptor.

[0711] As an alternative approach for receptor identification, the labeled polypeptides can be photoaffinity linked with cell membrane or extract preparations that express the receptor molecule. Cross-linked material is resolved by PAGE analysis and exposed to X-ray film. The labeled complex containing the receptors of the polypeptides can be excised, resolved into peptide fragments, and subjected to protein microsequencing. The amino acid sequence obtained from microsequencing would be used to design a set of degenerate oligonucleotide probes to screen a cDNA library to identify the genes encoding the putative receptors.

[0712] Moreover, the techniques of gene-shuffling, motif-shuffling, exon-shuffling, and/or codon-shuffling (collectively referred to as "DNA shuffling") may be employed to modulate the activities of the polypeptide of the present invention thereby effectively generating agonists and antagonists of the polypeptide of the present invention. See generally, U.S. Patent Nos. 5,605,793, 5,811,238, 5,830,721, 5,834,252, and 5,837,458, and Patten, P. A., *et al.*, *Curr. Opinion Biotechnol.* 8:724-33 (1997); Harayama, S. *Trends Biotechnol.* 16(2):76-82 (1998); Hansson, L. O., *et al.*, *J. Mol. Biol.* 287:265-76 (1999); and Lorenzo, M. M. and Blasco, R. *Biotechniques* 24(2):308-13 (1998); each of these patents and publications are hereby incorporated by reference). In one embodiment, alteration of polynucleotides and corresponding polypeptides may be achieved by DNA shuffling. DNA shuffling involves the assembly of two or more DNA segments into a desired molecule by homologous, or site-specific, recombination. In another embodiment, polynucleotides and corresponding polypeptides may be altered by being subjected to random mutagenesis by error-prone PCR, random nucleotide insertion or other methods prior to recombination. In another embodiment, one or more components, motifs, sections, parts, domains, fragments, etc., of the polypeptide of the present invention may

functional mimetic. (See, Coligan et al., Current Protocols in Immunology 1(2):Chapter 5 (1991)). Similarly, the molecule can be closely related to the natural receptor to which the polypeptide binds, or at least, a fragment of the receptor capable of being bound by the polypeptide (e.g., active site). In either case, the molecule can be rationally designed using known techniques.

[0705] Preferably, the screening for these molecules involves producing appropriate cells which express the polypeptide. Preferred cells include cells from mammals, yeast, *Drosophila*, or *E. coli*. Cells expressing the polypeptide (or cell membrane containing the expressed polypeptide) are then preferably contacted with a test compound potentially containing the molecule to observe binding, stimulation, or inhibition of activity of either the polypeptide or the molecule.

[0706] The assay may simply test binding of a candidate compound to the polypeptide, wherein binding is detected by a label, or in an assay involving competition with a labeled competitor. Further, the assay may test whether the candidate compound results in a signal generated by binding to the polypeptide.

[0707] Alternatively, the assay can be carried out using cell-free preparations, polypeptide/molecule affixed to a solid support, chemical libraries, or natural product mixtures. The assay may also simply comprise the steps of mixing a candidate compound with a solution containing a polypeptide, measuring polypeptide/molecule activity or binding, and comparing the polypeptide/molecule activity or binding to a standard.

[0708] Preferably, an ELISA assay can measure polypeptide level or activity in a sample (e.g., biological sample) using a monoclonal or polyclonal antibody. The antibody can measure polypeptide level or activity by either binding, directly or indirectly, to the polypeptide or by competing with the polypeptide for a substrate.

[0709] Additionally, the receptor to which the polypeptide of the present invention binds can be identified by numerous methods known to those of skill in the art, for example, ligand panning and FACS sorting (Coligan, et al., Current Protocols in Immun., 1(2), Chapter 5, (1991)). For example, expression cloning is employed wherein polyadenylated RNA is prepared from a cell responsive to the polypeptides, for example, NIH3T3 cells which are known to contain multiple receptors for the FGF family proteins, and SC-3 cells, and a cDNA library created from this RNA is divided into pools and used to transfect COS cells or other cells that are not responsive to the polypeptides.

hernia, umbilical hernia, ventral hernia), and intestinal diseases (e.g., cecal diseases (appendicitis, cecal neoplasms)).

Chemotaxis

[0700] Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention may have chemotaxis activity. A chemotactic molecule attracts or mobilizes cells (e.g., monocytes, fibroblasts, neutrophils, T-cells, mast cells, eosinophils, epithelial and/or endothelial cells) to a particular site in the body, such as inflammation, infection, or site of hyperproliferation. The mobilized cells can then fight off and/or heal the particular trauma or abnormality.

[0701] Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention may increase chemotactic activity of particular cells. These chemotactic molecules can then be used to treat inflammation, infection, hyperproliferative disorders, or any immune system disorder by increasing the number of cells targeted to a particular location in the body. For example, chemotactic molecules can be used to treat wounds and other trauma to tissues by attracting immune cells to the injured location. Chemotactic molecules of the present invention can also attract fibroblasts, which can be used to treat wounds.

[0702] It is also contemplated that polynucleotides or polypeptides, as well as agonists or antagonists of the present invention may inhibit chemotactic activity. These molecules could also be used to treat disorders. Thus, polynucleotides or polypeptides, as well as agonists or antagonists of the present invention could be used as an inhibitor of chemotaxis.

Binding Activity

[0703] A polypeptide of the present invention may be used to screen for molecules that bind to the polypeptide or for molecules to which the polypeptide binds. The binding of the polypeptide and the molecule may activate (agonist), increase, inhibit (antagonist), or decrease activity of the polypeptide or the molecule bound. Examples of such molecules include antibodies, oligonucleotides, proteins (e.g., receptors), or small molecules.

[0704] Preferably, the molecule is closely related to the natural ligand of the polypeptide, e.g., a fragment of the ligand, or a natural substrate, a ligand, a structural or

intussusception), intestinal perforation, intestinal polyps (colonic polyps, gardner syndrome, peutz-jeghers syndrome), jejunal diseases (jejunal neoplasms), malabsorption syndromes (blind loop syndrome, celiac disease, lactose intolerance, short bowel syndrome, tropical sprue, whipple's disease), mesenteric vascular occlusion, pneumatosis cystoides intestinalis, protein-losing enteropathies (intestinal lymphangiectasis), rectal diseases (anus diseases, fecal incontinence, hemorrhoids, proctitis, rectal fistula, rectal prolapse, rectocele), peptic ulcer (duodenal ulcer, peptic esophagitis, hemorrhage, perforation, stomach ulcer, Zollinger-Ellison syndrome), postgastrectomy syndromes (dumping syndrome), stomach diseases (e.g., achlorhydria, duodenogastric reflux (bile reflux), gastric antral vascular ectasia, gastric fistula, gastric outlet obstruction, gastritis (atrophic or hypertrophic), gastroparesis, stomach dilatation, stomach diverticulum, stomach neoplasms (gastric cancer, gastric polyps, gastric adenocarcinoma, hyperplastic gastric polyp), stomach rupture, stomach ulcer, stomach volvulus), tuberculosis, visceroptosis, vomiting (e.g., hematemesis, hyperemesis gravidarum, postoperative nausea and vomiting) and hemorrhagic colitis.

[0699] Further diseases and/or disorders of the gastrointestinal system include biliary tract diseases, such as, gastroschisis, fistula (e.g., biliary fistula, esophageal fistula, gastric fistula, intestinal fistula, pancreatic fistula), neoplasms (e.g., biliary tract neoplasms, esophageal neoplasms, such as adenocarcinoma of the esophagus, esophageal squamous cell carcinoma, gastrointestinal neoplasms, pancreatic neoplasms, such as adenocarcinoma of the pancreas, mucinous cystic neoplasm of the pancreas, pancreatic cystic neoplasms, pancreatoblastoma, and peritoneal neoplasms), esophageal disease (e.g., bullous diseases, candidiasis, glycogenic acanthosis, ulceration, barrett esophagus varices, atresia, cyst, diverticulum (e.g., Zenker's diverticulum), fistula (e.g., tracheoesophageal fistula), motility disorders (e.g., CREST syndrome, deglutition disorders, achalasia, spasm, gastroesophageal reflux), neoplasms, perforation (e.g., Boerhaave syndrome, Mallory-Weiss syndrome), stenosis, esophagitis, diaphragmatic hernia (e.g., hiatal hernia); gastrointestinal diseases, such as, gastroenteritis (e.g., cholera morbus, norwalk virus infection), hemorrhage (e.g., hematemesis, melena, peptic ulcer hemorrhage), stomach neoplasms (gastric cancer, gastric polyps, gastric adenocarcinoma, stomach cancer)), hernia (e.g., congenital diaphragmatic hernia, femoral hernia, inguinal hernia, obturator

Nodular regenerative hyperplasia)], malignant liver tumors [hepatocellular, hepatoblastoma, hepatocellular carcinoma, cholangiocellular, cholangiocarcinoma, cystadenocarcinoma, tumors of blood vessels, angiosarcoma, Kaposi's sarcoma, hemangioendothelioma, other tumors, embryonal sarcoma, fibrosarcoma, leiomyosarcoma, rhabdomyosarcoma, carcinosarcoma, teratoma, carcinoid, squamous carcinoma, primary lymphoma)], peliosis hepatis, erythrohepatic porphyria, hepatic porphyria (acute intermittent porphyria, porphyria cutanea tarda), Zellweger syndrome).

[0696] Pancreatic diseases and/or disorders include acute pancreatitis, chronic pancreatitis (acute necrotizing pancreatitis, alcoholic pancreatitis), neoplasms (adenocarcinoma of the pancreas, cystadenocarcinoma, insulinoma, gastrinoma, and glucagonoma, cystic neoplasms, islet-cell tumors, pancreoblastoma), and other pancreatic diseases (e.g., cystic fibrosis, cyst (pancreatic pseudocyst, pancreatic fistula, insufficiency)).

[0697] Gallbladder diseases include gallstones (cholelithiasis and choledocholithiasis), postcholecystectomy syndrome, diverticulosis of the gallbladder, acute cholecystitis, chronic cholecystitis, bile duct tumors, and mucocele.

[0698] Diseases and/or disorders of the large intestine include antibiotic-associated colitis, diverticulitis, ulcerative colitis, acquired megacolon, abscesses, fungal and bacterial infections, anorectal disorders (e.g., fissures, hemorrhoids), colonic diseases (colitis, colonic neoplasms [colon cancer, adenomatous colon polyps (e.g., villous adenoma), colon carcinoma, colorectal cancer], colonic diverticulitis, colonic diverticulosis, megacolon [Hirschsprung disease, toxic megacolon]; sigmoid diseases [proctocolitis, sigmoid neoplasms]), constipation, Crohn's disease, diarrhea (infantile diarrhea, dysentery), duodenal diseases (duodenal neoplasms, duodenal obstruction, duodenal ulcer, duodenitis), enteritis (enterocolitis), HIV enteropathy, ileal diseases (ileal neoplasms, ileitis), immunoproliferative small intestinal disease, inflammatory bowel disease (ulcerative colitis, Crohn's disease), intestinal atresia, parasitic diseases (anisakiasis, balantidiasis, blastocystis infections, cryptosporidiosis, dientamoebiasis, amebic dysentery, giardiasis), intestinal fistula (rectal fistula), intestinal neoplasms (cecal neoplasms, colonic neoplasms, duodenal neoplasms, ileal neoplasms, intestinal polyps, jejunal neoplasms, rectal neoplasms), intestinal obstruction (afferent loop syndrome, duodenal obstruction, impacted feces, intestinal pseudo-obstruction [cecal volvulus],

intolerance, celiac disease, duodenal ulcers, duodenitis, tropical sprue, Whipple's disease, intestinal lymphangiectasia, Crohn's disease, appendicitis, obstructions of the ileum, Meckel's diverticulum, multiple diverticula, failure of complete rotation of the small and large intestine, lymphoma, and bacterial and parasitic diseases (such as Traveler's diarrhea, typhoid and paratyphoid, cholera, infection by Roundworms (*Ascariasis lumbricoides*), Hookworms (*Ancylostoma duodenale*), Threadworms (*Enterobius vermicularis*), Tapeworms (*Taenia saginata*, *Echinococcus granulosus*, *Diphyllobothrium spp.*, and *T. solium*).

[0695] Liver diseases and/or disorders include intrahepatic cholestasis (alagille syndrome, biliary liver cirrhosis), fatty liver (alcoholic fatty liver, reye syndrome), hepatic vein thrombosis, hepatolenticular degeneration, hepatomegaly, hepatopulmonary syndrome, hepatorenal syndrome, portal hypertension (esophageal and gastric varices), liver abscess (amebic liver abscess), liver cirrhosis (alcoholic, biliary and experimental), alcoholic liver diseases (fatty liver, hepatitis, cirrhosis), parasitic (hepatic echinococcosis, fascioliasis, amebic liver abscess), jaundice (hemolytic, hepatocellular, and cholestatic), cholestasis, portal hypertension, liver enlargement, ascites, hepatitis (alcoholic hepatitis, animal hepatitis, chronic hepatitis (autoimmune, hepatitis B, hepatitis C, hepatitis D, drug induced), toxic hepatitis, viral human hepatitis (hepatitis A, hepatitis B, hepatitis C, hepatitis D, hepatitis E), Wilson's disease, granulomatous hepatitis, secondary biliary cirrhosis, hepatic encephalopathy, portal hypertension, varices, hepatic encephalopathy, primary biliary cirrhosis, primary sclerosing cholangitis, hepatocellular adenoma, hemangiomas, bile stones, liver failure (hepatic encephalopathy, acute liver failure), and liver neoplasms (angiomyolipoma, calcified liver metastases, cystic liver metastases, epithelial tumors, fibrolamellar hepatocarcinoma, focal nodular hyperplasia, hepatic adenoma, hepatobiliary cystadenoma, hepatoblastoma, hepatocellular carcinoma, hepatoma, liver cancer, liver hemangioendothelioma, mesenchymal hamartoma, mesenchymal tumors of liver, nodular regenerative hyperplasia, benign liver tumors (Hepatic cysts [Simple cysts, Polycystic liver disease, Hepatobiliary cystadenoma, Choledochal cyst], Mesenchymal tumors [Mesenchymal hamartoma, Infantile hemangioendothelioma, Hemangioma, Peliosis hepatis, Lipomas, Inflammatory pseudotumor, Miscellaneous], Epithelial tumors [Bile duct epithelium (Bile duct hamartoma, Bile duct adenoma), Hepatocyte (Adenoma, Focal nodular hyperplasia,

includes pressure ulcers, ulcers associated with vascular insufficiency, surgical, and traumatic wounds.

[0691] Similarly, nerve and brain tissue could also be regenerated by using polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, to proliferate and differentiate nerve cells. Diseases that could be treated using this method include central and peripheral nervous system diseases, neuropathies, or mechanical and traumatic disorders (e.g., spinal cord disorders, head trauma, cerebrovascular disease, and stroke). Specifically, diseases associated with peripheral nerve injuries, peripheral neuropathy (e.g., resulting from chemotherapy or other medical therapies), localized neuropathies, and central nervous system diseases (e.g., Alzheimer's disease, Parkinson's disease, Huntington's disease, amyotrophic lateral sclerosis, and Shy-Drager syndrome), could all be treated using the polynucleotides or polypeptides, as well as agonists or antagonists of the present invention.

Gastrointestinal Disorders

[0692] Polynucleotides or polypeptides, or agonists or antagonists of the present invention, may be used to treat, prevent, diagnose, and/or prognose gastrointestinal disorders, including inflammatory diseases and/or conditions, infections, cancers (e.g., intestinal neoplasms (carcinoid tumor of the small intestine, non-Hodgkin's lymphoma of the small intestine, small bowel lymphoma)), and ulcers, such as peptic ulcers.

[0693] Gastrointestinal disorders include dysphagia, odynophagia, inflammation of the esophagus, peptic esophagitis, gastric reflux, submucosal fibrosis and stricturing, Mallory-Weiss lesions, leiomyomas, lipomas, epidermal cancers, adeoncarcinomas, gastric retention disorders, gastroenteritis, gastric atrophy, gastric/stomach cancers, polyps of the stomach, autoimmune disorders such as pernicious anemia, pyloric stenosis, gastritis (bacterial, viral, eosinophilic, stress-induced, chronic erosive, atrophic, plasma cell, and Ménétrier's), and peritoneal diseases (e.g., chyloperitoneum, hemoperitoneum, mesenteric cyst, mesenteric lymphadenitis, mesenteric vascular occlusion, panniculitis, neoplasms, peritonitis, pneumoperitoneum, bubphrenic abscess,).

[0694] Gastrointestinal disorders also include disorders associated with the small intestine, such as malabsorption syndromes, distension, irritable bowel syndrome, sugar

complications, and toxoplasmosis. polynucleotides or polypeptides, or agonists or antagonists of the invention, can be used to treat, prevent, and/or diagnose any of these symptoms or diseases. In specific embodiments, polynucleotides, polypeptides, or agonists or antagonists of the invention are used to treat, prevent, and/or diagnose malaria.

[0687] Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention could either be by administering an effective amount of a polypeptide to the patient, or by removing cells from the patient, supplying the cells with a polynucleotide of the present invention, and returning the engineered cells to the patient (ex vivo therapy). Moreover, the polypeptide or polynucleotide of the present invention can be used as an antigen in a vaccine to raise an immune response against infectious disease.

Regeneration

[0688] Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention can be used to differentiate, proliferate, and attract cells, leading to the regeneration of tissues. (See, Science 276:59-87 (1997)). The regeneration of tissues could be used to repair, replace, or protect tissue damaged by congenital defects, trauma (wounds, burns, incisions, or ulcers), age, disease (e.g. osteoporosis, osteoarthritis, periodontal disease, liver failure), surgery, including cosmetic plastic surgery, fibrosis, reperfusion injury, or systemic cytokine damage.

[0689] Tissues that could be regenerated using the present invention include organs (e.g., pancreas, liver, intestine, kidney, skin, endothelium), muscle (smooth, skeletal or cardiac), vasculature (including vascular and lymphatics), nervous, hematopoietic, and skeletal (bone, cartilage, tendon, and ligament) tissue. Preferably, regeneration occurs without or decreased scarring. Regeneration also may include angiogenesis.

[0690] Moreover, polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, may increase regeneration of tissues difficult to heal. For example, increased tendon/ligament regeneration would quicken recovery time after damage. Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention could also be used prophylactically in an effort to avoid damage. Specific diseases that could be treated include of tendinitis, carpal tunnel syndrome, and other tendon or ligament defects. A further example of tissue regeneration of non-healing wounds

Treponema spp., Leptospira spp., Borrelia spp.), Shigella spp., Staphylococcus (e.g., *Staphylococcus aureus*), Meningioccus, Pneumococcus and Streptococcus (e.g., *Streptococcus pneumoniae* and Groups A, B, and C Streptococci), and Ureaplasmas. These bacterial, parasitic, and fungal families can cause diseases or symptoms, including, but not limited to: antibiotic-resistant infections, bacteremia, endocarditis, septicemia, eye infections (e.g., conjunctivitis), uveitis, tuberculosis, gingivitis, bacterial diarrhea, opportunistic infections (e.g., AIDS related infections), paronychia, prosthesis-related infections, dental caries, Reiter's Disease, respiratory tract infections, such as Whooping Cough or Empyema, sepsis, Lyme Disease, Cat-Scratch Disease, dysentery, paratyphoid fever, food poisoning, Legionella disease, chronic and acute inflammation, erythema, yeast infections, typhoid, pneumonia, gonorrhea, meningitis (e.g., meningitis types A and B), chlamydia, syphilis, diphtheria, leprosy, brucellosis, peptic ulcers, anthrax, spontaneous abortions, birth defects, pneumonia, lung infections, ear infections, deafness, blindness, lethargy, malaise, vomiting, chronic diarrhea, Crohn's disease, colitis, vaginosis, sterility, pelvic inflammatory diseases, candidiasis, paratuberculosis, tuberculosis, lupus, botulism, gangrene, tetanus, impetigo, Rheumatic Fever, Scarlet Fever, sexually transmitted diseases, skin diseases (e.g., cellulitis, dermatocycoses), toxemia, urinary tract infections, wound infections, noscomial infections. Polynucleotides or polypeptides, agonists or antagonists of the invention, can be used to treat or detect any of these symptoms or diseases. In specific embodiments, polynucleotides, polypeptides, agonists or antagonists of the invention are used to treat: tetanus, diphtheria, botulism, and/or meningitis type B.

[0686] Moreover, parasitic agents causing disease or symptoms that can be treated, prevented, and/or diagnosed by a polynucleotide or polypeptide and/or agonist or antagonist of the present invention include, but not limited to, the following families or class: Amebiasis, Babesiosis, Coccidiosis, Cryptosporidiosis, Dientamoebiasis, Dourine, Ectoparasitic, Giardiasis, Helminthiasis, Leishmaniasis, Schistosoma, Theileriasis, Toxoplasmosis, Trypanosomiasis, and Trichomonas and Sporozoans (e.g., *Plasmodium virax*, *Plasmodium falciparum*, *Plasmodium malariae* and *Plasmodium ovale*). These parasites can cause a variety of diseases or symptoms, including, but not limited to: Scabies, Trombiculiasis, eye infections, intestinal disease (e.g., dysentery, giardiasis), liver disease, lung disease, opportunistic infections (e.g., AIDS related), malaria, pregnancy

bronchiollitis, respiratory syncytial virus, encephalitis, eye infections (e.g., conjunctivitis, keratitis), chronic fatigue syndrome, hepatitis (A, B, C, E, Chronic Active, Delta), Japanese B encephalitis, Junin, Chikungunya, Rift Valley fever, yellow fever, meningitis, opportunistic infections (e.g., AIDS), pneumonia, Burkitt's Lymphoma, chickenpox, hemorrhagic fever, Measles, Mumps, Parainfluenza, Rabies, the common cold, Polio, leukemia, Rubella, sexually transmitted diseases, skin diseases (e.g., Kaposi's, warts), and viremia. polynucleotides or polypeptides, or agonists or antagonists of the invention, can be used to treat or detect any of these symptoms or diseases. In specific embodiments, polynucleotides, polypeptides, or agonists or antagonists of the invention are used to treat: meningitis, Dengue, EBV, and/or hepatitis (e.g., hepatitis B). In an additional specific embodiment polynucleotides, polypeptides, or agonists or antagonists of the invention are used to treat patients nonresponsive to one or more other commercially available hepatitis vaccines. In a further specific embodiment polynucleotides, polypeptides, or agonists or antagonists of the invention are used to treat AIDS.

[0685] Similarly, bacterial and fungal agents that can cause disease or symptoms and that can be treated or detected by a polynucleotide or polypeptide and/or agonist or antagonist of the present invention include, but not limited to, the following Gram-Negative and Gram-positive bacteria, bacterial families, and fungi: Actinomyces (e.g., Norcardia), Acinetobacter, *Cryptococcus neoformans*, Aspergillus, Bacillaceae (e.g., *Bacillus anthracis*), Bacteroides (e.g., *Bacteroides fragilis*), Blastomycosis, Bordetella, Borrelia (e.g., *Borrelia burgdorferi*), Brucella, Candidia, Campylobacter, Chlamydia, Clostridium (e.g., *Clostridium botulinum*, *Clostridium difficile*, *Clostridium perfringens*, *Clostridium tetani*), Coccidioides, Corynebacterium (e.g., *Corynebacterium diphtheriae*), Cryptococcus, Dermatocycoses, *E. coli* (e.g., Enterotoxigenic *E. coli* and Enterohemorrhagic *E. coli*), Enterobacter (e.g. *Enterobacter aerogenes*), Enterobacteriaceae (Klebsiella, Salmonella (e.g., *Salmonella typhi*, *Salmonella enteritidis*, *Salmonella typhi*), Serratia, Yersinia, Shigella), Erysipelothrix, Haemophilus (e.g., *Haemophilus influenza* type B), Helicobacter, Legionella (e.g., *Legionella pneumophila*), Leptospira, Listeria (e.g., *Listeria monocytogenes*), Mycoplasma, Mycobacterium (e.g., *Mycobacterium leprae* and *Mycobacterium tuberculosis*), Vibrio (e.g., *Vibrio cholerae*), Neisseriaceae (e.g., *Neisseria gonorrhea*, *Neisseria meningitidis*), Pasteurellaceae, Proteus, Pseudomonas (e.g., *Pseudomonas aeruginosa*), Rickettsiaceae, Spirochetes (e.g.,

embolism, endotoxemia, pyelonephritis, saphenous thrombophlebitis, mastitis, cystitis, postpartum hemorrhage, and inverted uterus.

[0682] Other disorders and/or diseases of the female reproductive system that may be diagnosed, treated, and/or prevented by the polynucleotides, polypeptides, and agonists or antagonists of the present invention include, for example, Turner's syndrome, pseudohermaphroditism, premenstrual syndrome, pelvic inflammatory disease, pelvic congestion (vascular engorgement), frigidity, anorgasmia, dyspareunia, ruptured fallopian tube, and Mittelschmerz.

Infectious Disease

[0683] Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention can be used to treat or detect infectious agents. For example, by increasing the immune response, particularly increasing the proliferation and differentiation of B and/or T cells, infectious diseases may be treated. The immune response may be increased by either enhancing an existing immune response, or by initiating a new immune response. Alternatively, polynucleotides or polypeptides, as well as agonists or antagonists of the present invention may also directly inhibit the infectious agent, without necessarily eliciting an immune response.

[0684] Viruses are one example of an infectious agent that can cause disease or symptoms that can be treated or detected by a polynucleotide or polypeptide and/or agonist or antagonist of the present invention. Examples of viruses, include, but are not limited to Examples of viruses, include, but are not limited to the following DNA and RNA viruses and viral families: Arbovirus, Adenoviridae, Arenaviridae, Arterivirus, Birnaviridae, Bunyaviridae, Caliciviridae, Circoviridae, Coronaviridae, Dengue, EBV, HIV, Flaviviridae, Hepadnaviridae (Hepatitis), Herpesviridae (such as, Cytomegalovirus, Herpes Simplex, Herpes Zoster), Mononegavirus (e.g., Paramyxoviridae, Morbillivirus, Rhabdoviridae), Orthomyxoviridae (e.g., Influenza A, Influenza B, and parainfluenza), Papilloma virus, Papovaviridae, Parvoviridae, Picornaviridae, Poxviridae (such as Smallpox or Vaccinia), Reoviridae (e.g., Rotavirus), Retroviridae (HTLV-I, HTLV-II, Lentivirus), and Togaviridae (e.g., Rubivirus). Viruses falling within these families can cause a variety of diseases or symptoms, including, but not limited to: arthritis,

of the ovary, ovarian papillary serous adenocarcinoma, ovarian mucinous adenocarcinoma, and Ovarian Krukenberg tumors).

[0678] Cervical diseases and/or disorders include cervicitis, chronic cervicitis, mucopurulent cervicitis, cervical dysplasia, cervical polyps, Nabothian cysts, cervical erosion, cervical incompetence, and cervical neoplasms (including, for example, cervical carcinoma, squamous metaplasia, squamous cell carcinoma, adenosquamous cell neoplasia, and columnar cell neoplasia).

[0679] Additionally, diseases and/or disorders of the reproductive system include disorders and/or diseases of pregnancy, including miscarriage and stillbirth, such as early abortion, late abortion, spontaneous abortion, induced abortion, therapeutic abortion, threatened abortion, missed abortion, incomplete abortion, complete abortion, habitual abortion, missed abortion, and septic abortion; ectopic pregnancy, anemia, Rh incompatibility, vaginal bleeding during pregnancy, gestational diabetes, intrauterine growth retardation, polyhydramnios, HELLP syndrome, abruptio placentae, placenta previa, hyperemesis, preeclampsia, eclampsia, herpes gestationis, and urticaria of pregnancy. Additionally, the polynucleotides, polypeptides, and agonists or antagonists of the present invention may be used in the diagnosis, treatment, and/or prevention of diseases that can complicate pregnancy, including heart disease, heart failure, rheumatic heart disease, congenital heart disease, mitral valve prolapse, high blood pressure, anemia, kidney disease, infectious disease (e.g., rubella, cytomegalovirus, toxoplasmosis, infectious hepatitis, chlamydia, HIV, AIDS, and genital herpes), diabetes mellitus, Graves' disease, thyroiditis, hypothyroidism, Hashimoto's thyroiditis, chronic active hepatitis, cirrhosis of the liver, primary biliary cirrhosis, asthma, systemic lupus erythematosus, rheumatoid arthritis, myasthenia gravis, idiopathic thrombocytopenic purpura, appendicitis, ovarian cysts, gallbladder disorders, and obstruction of the intestine.

[0680] Complications associated with labor and parturition include premature rupture of the membranes, pre-term labor, post-term pregnancy, postmaturity, labor that progresses too slowly, fetal distress (e.g., abnormal heart rate (fetal or maternal), breathing problems, and abnormal fetal position), shoulder dystocia, prolapsed umbilical cord, amniotic fluid embolism, and aberrant uterine bleeding.

[0681] Further, diseases and/or disorders of the postdelivery period, including endometritis, myometritis, parametritis, peritonitis, pelvic thrombophlebitis, pulmonary

[0674] Other disorders and/or diseases of the male reproductive system include, for example, Klinefelter's syndrome, Young's syndrome, premature ejaculation, diabetes mellitus, cystic fibrosis, Kartagener's syndrome, high fever, multiple sclerosis, and gynecomastia.

[0675] Further, the polynucleotides, polypeptides, and agonists or antagonists of the present invention may be used in the diagnosis, treatment, and/or prevention of diseases and/or disorders of the vagina and vulva, including bacterial vaginosis, candida vaginitis, herpes simplex virus, chancroid, granuloma inguinale, lymphogranuloma venereum, scabies, human papillomavirus, vaginal trauma, vulvar trauma, adenosis, chlamydia vaginitis, gonorrhea, trichomonas vaginitis, condyloma acuminatum, syphilis, molluscum contagiosum, atrophic vaginitis, Paget's disease, lichen sclerosus, lichen planus, vulvodynia, toxic shock syndrome, vaginismus, vulvovaginitis, vulvar vestibulitis, and neoplastic disorders, such as squamous cell hyperplasia, clear cell carcinoma, basal cell carcinoma, melanomas, cancer of Bartholin's gland, and vulvar intraepithelial neoplasia.

[0676] Disorders and/or diseases of the uterus include dysmenorrhea, retroverted uterus, endometriosis, fibroids, adenomyosis, anovulatory bleeding, amenorrhea, Cushing's syndrome, hydatidiform moles, Asherman's syndrome, premature menopause, precocious puberty, uterine polyps, dysfunctional uterine bleeding (e.g., due to aberrant hormonal signals), and neoplastic disorders, such as adenocarcinomas, leiomyosarcomas, and sarcomas. Additionally, the polypeptides, polynucleotides, or agonists or antagonists of the invention may be useful as a marker or detector of, as well as in the diagnosis, treatment, and/or prevention of congenital uterine abnormalities, such as bicornuate uterus, septate uterus, simple unicornuate uterus, unicornuate uterus with a noncavitary rudimentary horn, unicornuate uterus with a non-communicating cavitary rudimentary horn, unicornuate uterus with a communicating cavitary horn, arcuate uterus, uterine didelphys, and T-shaped uterus.

[0677] Ovarian diseases and/or disorders include anovulation, polycystic ovary syndrome (Stein-Leventhal syndrome), ovarian cysts, ovarian hypofunction, ovarian insensitivity to gonadotropins, ovarian overproduction of androgens, right ovarian vein syndrome, amenorrhea, hirsutism, and ovarian cancer (including, but not limited to, primary and secondary cancerous growth, Sertoli-Leydig tumors, endometrioid carcinoma

and bilateral), anorchia, ectopic testis, epididymitis and orchitis (typically resulting from infections such as, for example, gonorrhea, mumps, tuberculosis, and syphilis), testicular torsion, vasitis nodosa, germ cell tumors (e.g., seminomas, embryonal cell carcinomas, teratocarcinomas, choriocarcinomas, yolk sac tumors, and teratomas), stromal tumors (e.g., Leydig cell tumors), hydrocele, hematocele, varicocele, spermatocele, inguinal hernia, and disorders of sperm production (e.g., immotile cilia syndrome, aspermia, asthenozoospermia, azoospermia, oligospermia, and teratozoospermia).

[0671] Reproductive system disorders also include disorders of the prostate gland, such as acute non-bacterial prostatitis, chronic non-bacterial prostatitis, acute bacterial prostatitis, chronic bacterial prostatitis, prostatodystonia, prostatosis, granulomatous prostatitis, malacoplakia, benign prostatic hypertrophy or hyperplasia, and prostate neoplastic disorders, including adenocarcinomas, transitional cell carcinomas, ductal carcinomas, and squamous cell carcinomas.

[0672] Additionally, the compositions of the invention may be useful in the diagnosis, treatment, and/or prevention of disorders or diseases of the penis and urethra, including inflammatory disorders, such as balanoposthitis, balanitis xerotica obliterans, phimosis, paraphimosis, syphilis, herpes simplex virus, gonorrhea, non-gonococcal urethritis, chlamydia, mycoplasma, trichomonas, HIV, AIDS, Reiter's syndrome, condyloma acuminatum, condyloma latum, and pearly penile papules; urethral abnormalities, such as hypospadias, epispadias, and phimosis; premalignant lesions, including Erythroplasia of Queyrat, Bowen's disease, Bowenoid papulosis, giant condyloma of Buscke-Lowenstein, and verrucous carcinoma; penile cancers, including squamous cell carcinomas, carcinoma in situ, verrucous carcinoma, and disseminated penile carcinoma; urethral neoplastic disorders, including penile urethral carcinoma, bulbomembranous urethral carcinoma, and prostatic urethral carcinoma; and erectile disorders, such as priapism, Peyronie's disease, erectile dysfunction, and impotence.

[0673] Moreover, diseases and/or disorders of the vas deferens include vasculitis and CBAVD (congenital bilateral absence of the vas deferens); additionally, the polynucleotides, polypeptides, and agonists or antagonists of the present invention may be used in the diagnosis, treatment, and/or prevention of diseases and/or disorders of the seminal vesicles, including hydatid disease, congenital chloride diarrhea, and polycystic kidney disease.

coupling defect, thymic aplasia, Hurthle cell tumours of the thyroid, thyroid cancer, thyroid carcinoma, Medullary thyroid carcinoma; disorders and/or diseases of the parathyroid, such as, for example, hyperparathyroidism, hypoparathyroidism; disorders and/or diseases of the hypothalamus.

[0666] In addition, endocrine system and/or hormone imbalance disorders and/or diseases may also include disorders and/or diseases of the testes or ovaries, including cancer. Other disorders and/or diseases of the testes or ovaries further include, for example, ovarian cancer, polycystic ovary syndrome, Klinefelter's syndrome, vanishing testes syndrome (bilateral anorchia), congenital absence of Leydig's cells, cryptorchidism, Noonan's syndrome, myotonic dystrophy, capillary haemangioma of the testis (benign), neoplasias of the testis and neo-testis.

[0667] Moreover, endocrine system and/or hormone imbalance disorders and/or diseases may also include disorders and/or diseases such as, for example, polyglandular deficiency syndromes, pheochromocytoma, neuroblastoma, multiple Endocrine neoplasia, and disorders and/or cancers of endocrine tissues.

[0668] In another embodiment, a polypeptide of the invention, or polynucleotides, antibodies, agonists, or antagonists corresponding to that polypeptide, may be used to diagnose, prognose, prevent, and/or treat endocrine diseases and/or disorders associated with the tissue(s) in which the polypeptide of the invention is expressed, including one, two, three, four, five, or more tissues disclosed in Table 1, column 8 (Tissue Distribution Library Code).

Reproductive System Disorders

[0669] The polynucleotides or polypeptides, or agonists or antagonists of the invention may be used for the diagnosis, treatment, or prevention of diseases and/or disorders of the reproductive system. Reproductive system disorders that can be treated by the compositions of the invention, include, but are not limited to, reproductive system injuries, infections, neoplastic disorders, congenital defects, and diseases or disorders which result in infertility, complications with pregnancy, labor, or parturition, and postpartum difficulties.

[0670] Reproductive system disorders and/or diseases include diseases and/or disorders of the testes, including testicular atrophy, testicular feminization, cryptorchism (unilateral

Endocrine Disorders

[0662] Polynucleotides or polypeptides, or agonists or antagonists of the present invention, may be used to treat, prevent, diagnose, and/or prognose disorders and/or diseases related to hormone imbalance, and/or disorders or diseases of the endocrine system.

[0663] Hormones secreted by the glands of the endocrine system control physical growth, sexual function, metabolism, and other functions. Disorders may be classified in two ways: disturbances in the production of hormones, and the inability of tissues to respond to hormones. The etiology of these hormone imbalance or endocrine system diseases, disorders or conditions may be genetic, somatic, such as cancer and some autoimmune diseases, acquired (e.g., by chemotherapy, injury or toxins), or infectious. Moreover, polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention can be used as a marker or detector of a particular disease or disorder related to the endocrine system and/or hormone imbalance.

[0664] Endocrine system and/or hormone imbalance and/or diseases encompass disorders of uterine motility including, but not limited to: complications with pregnancy and labor (e.g., pre-term labor, post-term pregnancy, spontaneous abortion, and slow or stopped labor); and disorders and/or diseases of the menstrual cycle (e.g., dysmenorrhea and endometriosis).

[0665] Endocrine system and/or hormone imbalance disorders and/or diseases include disorders and/or diseases of the pancreas, such as, for example, diabetes mellitus, diabetes insipidus, congenital pancreatic agenesis, pheochromocytoma--islet cell tumor syndrome; disorders and/or diseases of the adrenal glands such as, for example, Addison's Disease, corticosteroid deficiency, virilizing disease, hirsutism, Cushing's Syndrome, hyperaldosteronism, pheochromocytoma; disorders and/or diseases of the pituitary gland, such as, for example, hyperpituitarism, hypopituitarism, pituitary dwarfism, pituitary adenoma, panhypopituitarism, acromegaly, gigantism; disorders and/or diseases of the thyroid, including but not limited to, hyperthyroidism, hypothyroidism, Plummer's disease, Graves' disease (toxic diffuse goiter), toxic nodular goiter, thyroiditis (Hashimoto's thyroiditis, subacute granulomatous thyroiditis, and silent lymphocytic thyroiditis), Pendred's syndrome, myxedema, cretinism, thyrotoxicosis, thyroid hormone

spasm such as trismus, unconsciousness such as coma, persistent vegetative state and syncope and vertigo, neuromuscular diseases such as amyotonia congenita, amyotrophic lateral sclerosis, Lambert-Eaton Myasthenic Syndrome, motor neuron disease, muscular atrophy such as spinal muscular atrophy, Charcot-Marie Disease and Werdnig-Hoffmann Disease, Postpoliomyelitis Syndrome, Muscular Dystrophy, Myasthenia Gravis, Myotonia Atrophica, Myotonia Confenita, Nemaline Myopathy, Familial Periodic Paralysis, Multiplex Paramyoclonus, Tropical Spastic Paraparesis and Stiff-Man Syndrome, peripheral nervous system diseases such as acrodynia, amyloid neuropathies, autonomic nervous system diseases such as Adie's Syndrome, Barre-Lieou Syndrome, Familial Dysautonomia, Horner's Syndrome, Reflex Sympathetic Dystrophy and Shy-Drager Syndrome, Cranial Nerve Diseases such as Acoustic Nerve Diseases such as Acoustic Neuroma which includes Neurofibromatosis 2, Facial Nerve Diseases such as Facial Neuralgia, Melkersson-Rosenthal Syndrome, ocular motility disorders which includes amblyopia, nystagmus, oculomotor nerve paralysis, ophthalmoplegia such as Duane's Syndrome, Horner's Syndrome, Chronic Progressive External Ophthalmoplegia which includes Kearns Syndrome, Strabismus such as Esotropia and Exotropia, Oculomotor Nerve Paralysis, Optic Nerve Diseases such as Optic Atrophy which includes Hereditary Optic Atrophy, Optic Disk Drusen, Optic Neuritis such as Neuromyelitis Optica, Papilledema, Trigeminal Neuralgia, Vocal Cord Paralysis, Demyelinating Diseases such as Neuromyelitis Optica and Swayback, and Diabetic neuropathies such as diabetic foot.

[0661] Additional neurologic diseases which can be treated or detected with polynucleotides, polypeptides, agonists, and/or antagonists of the present invention include nerve compression syndromes such as carpal tunnel syndrome, tarsal tunnel syndrome, thoracic outlet syndrome such as cervical rib syndrome, ulnar nerve compression syndrome, neuralgia such as causalgia, cervico-brachial neuralgia, facial neuralgia and trigeminal neuralgia, neuritis such as experimental allergic neuritis, optic neuritis, polyneuritis, polyradiculoneuritis and radiculities such as polyradiculitis, hereditary motor and sensory neuropathies such as Charcot-Marie Disease, Hereditary Optic Atrophy, Refsum's Disease, Hereditary Spastic Paraplegia and Werdnig-Hoffmann Disease, Hereditary Sensory and Autonomic Neuropathies which include Congenital Analgesia and Familial Dysautonomia, POEMS Syndrome, Sciatica, Gustatory Sweating and Tetany).

oculocerebrorenal syndrome, phenylketonuria such as maternal phenylketonuria, Prader-Willi Syndrome, Rett Syndrome, Rubinstein-Taybi Syndrome, Tuberous Sclerosis, WAGR Syndrome, nervous system abnormalities such as holoprosencephaly, neural tube defects such as anencephaly which includes hydrangencephaly, Arnold-Chairi Deformity, encephalocele, meningocele, meningomyelocele, spinal dysraphism such as spina bifida cystica and spina bifida occulta.

[0660] Additional neurologic diseases which can be treated or detected with polynucleotides, polypeptides, agonists, and/or antagonists of the present invention include hereditary motor and sensory neuropathies which include Charcot-Marie Disease, Hereditary optic atrophy, Refsum's Disease, hereditary spastic paraplegia, Werdnig-Hoffmann Disease, Hereditary Sensory and Autonomic Neuropathies such as Congenital Analgesia and Familial Dysautonomia, Neurologic manifestations (such as agnosia that include Gerstmann's Syndrome, Amnesia such as retrograde amnesia, apraxia, neurogenic bladder, cataplexy, communicative disorders such as hearing disorders that includes deafness, partial hearing loss, loudness recruitment and tinnitus, language disorders such as aphasia which include agraphia, anomia, broca aphasia, and Wernicke Aphasia, Dyslexia such as Acquired Dyslexia, language development disorders, speech disorders such as aphasia which includes anomia, broca aphasia and Wernicke Aphasia, articulation disorders, communicative disorders such as speech disorders which include dysarthria, echolalia, mutism and stuttering, voice disorders such as aphonia and hoarseness, decerebrate state, delirium, fasciculation, hallucinations, meningism, movement disorders such as angelman syndrome, ataxia, athetosis, chorea, dystonia, hypokinesia, muscle hypotonia, myoclonus, tic, torticollis and tremor, muscle hypertonia such as muscle rigidity such as stiff-man syndrome, muscle spasticity, paralysis such as facial paralysis which includes Herpes Zoster Oticus, Gastroparesis, Hemiplegia, ophthalmoplegia such as diplopia, Duane's Syndrome, Horner's Syndrome, Chronic progressive external ophthalmoplegia such as Kearns Syndrome, Bulbar Paralysis, Tropical Spastic Paraparesis, Paraplegia such as Brown-Sequard Syndrome, quadriplegia, respiratory paralysis and vocal cord paralysis, paresis, phantom limb, taste disorders such as ageusia and dysgeusia, vision disorders such as amblyopia, blindness, color vision defects, diplopia, hemianopsia, scotoma and subnormal vision, sleep disorders such as hypersomnia which includes Kleine-Levin Syndrome, insomnia, and somnambulism,

[0658] Additional neurologic diseases which can be treated or detected with polynucleotides, polypeptides, agonists, and/or antagonists of the present invention include meningitis such as arachnoiditis, aseptic meningitis such as viral meningitis which includes lymphocytic choriomeningitis, Bacterial meningitis which includes Haemophilus Meningitis, Listeria Meningitis, Meningococcal Meningitis such as Waterhouse-Friderichsen Syndrome, Pneumococcal Meningitis and meningeal tuberculosis, fungal meningitis such as Cryptococcal Meningitis, subdural effusion, meningoencephalitis such as uvemeningoencephalitic syndrome, myelitis such as transverse myelitis, neurosyphilis such as tabes dorsalis, poliomyelitis which includes bulbar poliomyelitis and postpoliomyelitis syndrome, prion diseases (such as Creutzfeldt-Jakob Syndrome, Bovine Spongiform Encephalopathy, Gerstmann-Straussler Syndrome, Kuru, Scrapie), and cerebral toxoplasmosis.

[0659] Additional neurologic diseases which can be treated or detected with polynucleotides, polypeptides, agonists, and/or antagonists of the present invention include central nervous system neoplasms such as brain neoplasms that include cerebellar neoplasms such as infratentorial neoplasms, cerebral ventricle neoplasms such as choroid plexus neoplasms, hypothalamic neoplasms and supratentorial neoplasms, meningeal neoplasms, spinal cord neoplasms which include epidural neoplasms, demyelinating diseases such as Canavan Diseases, diffuse cerebral sclerolosis which includes adrenoleukodystrophy, encephalitis periaxialis, globoid cell leukodystrophy, diffuse cerebral sclerosis such as metachromatic leukodystrophy, allergic encephalomyelitis, necrotizing hemorrhagic encephalomyelitis, progressive multifocal leukoencephalopathy, multiple sclerosis, central pontine myelinolysis, transverse myelitis, neuromyelitis optica, Scrapie, Swayback, Chronic Fatigue Syndrome, Visna, High Pressure Nervous Syndrome, Meningism, spinal cord diseases such as amyotonia congenita, amyotrophic lateral sclerosis, spinal muscular atrophy such as Werdnig-Hoffmann Disease, spinal cord compression, spinal cord neoplasms such as epidural neoplasms, syringomyelia, Tabes Dorsalis, Stiff-Man Syndrome, mental retardation such as Angelman Syndrome, Cri-du-Chat Syndrome, De Lange's Syndrome, Down Syndrome, Gangliosidoses such as gangliosidoses G(M1), Sandhoff Disease, Tay-Sachs Disease, Hartnup Disease, homocystinuria, Laurence-Moon- Biedl Syndrome, Lesch-Nyhan Syndrome, Maple Syrup Urine Disease, mucopolipidosis such as fucosidosis, neuronal ceroid-lipofuscinosis,

malformations, cerebral artery diseases, cerebral embolism and thrombosis such as carotid artery thrombosis, sinus thrombosis and Wallenberg's Syndrome, cerebral hemorrhage such as epidural hematoma, subdural hematoma and subarachnoid hemorrhage, cerebral infarction, cerebral ischemia such as transient cerebral ischemia, Subclavian Steal Syndrome and vertebrobasilar insufficiency, vascular dementia such as multi-infarct dementia, periventricular leukomalacia, vascular headache such as cluster headache and migraine.

[0656] Additional neurologic diseases which can be treated or detected with polynucleotides, polypeptides, agonists, and/or antagonists of the present invention include dementia such as AIDS Dementia Complex, presenile dementia such as Alzheimer's Disease and Creutzfeldt-Jakob Syndrome, senile dementia such as Alzheimer's Disease and progressive supranuclear palsy, vascular dementia such as multi-infarct dementia, encephalitis which include encephalitis periaxialis, viral encephalitis such as epidemic encephalitis, Japanese Encephalitis, St. Louis Encephalitis, tick-borne encephalitis and West Nile Fever, acute disseminated encephalomyelitis, meningoencephalitis such as uveomeningoencephalitic syndrome, Postencephalitic Parkinson Disease and subacute sclerosing panencephalitis, encephalomalacia such as periventricular leukomalacia, epilepsy such as generalized epilepsy which includes infantile spasms, absence epilepsy, myoclonic epilepsy which includes MERRF Syndrome, tonic-clonic epilepsy, partial epilepsy such as complex partial epilepsy, frontal lobe epilepsy and temporal lobe epilepsy, post-traumatic epilepsy, status epilepticus such as Epilepsia Partialis Continua, and Hallervorden-Spatz Syndrome.

[0657] Additional neurologic diseases which can be treated or detected with polynucleotides, polypeptides, agonists, and/or antagonists of the present invention include hydrocephalus such as Dandy-Walker Syndrome and normal pressure hydrocephalus, hypothalamic diseases such as hypothalamic neoplasms, cerebral malaria, narcolepsy which includes cataplexy, bulbar poliomyelitis, cerebri pseudotumor, Rett Syndrome, Reye's Syndrome, thalamic diseases, cerebral toxoplasmosis, intracranial tuberculoma and Zellweger Syndrome, central nervous system infections such as AIDS Dementia Complex, Brain Abscess, subdural empyema, encephalomyelitis such as Equine Encephalomyelitis, Venezuelan Equine Encephalomyelitis, Necrotizing Hemorrhagic Encephalomyelitis, Visna, and cerebral malaria.

cerebral arteriovenous malformations, cerebral artery diseases, cerebral embolism and thrombosis (e.g., carotid artery thrombosis, sinus thrombosis, or Wallenberg's Syndrome), cerebral hemorrhage (e.g., epidural or subdural hematoma, or subarachnoid hemorrhage), cerebral infarction, cerebral ischemia (e.g., transient cerebral ischemia, Subclavian Steal Syndrome, or vertebrobasilar insufficiency), vascular dementia (e.g., multi-infarct), leukomalacia, periventricular, and vascular headache (e.g., cluster headache or migraines).

[0653] In accordance with yet a further aspect of the present invention, there is provided a process for utilizing polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, for therapeutic purposes, for example, to stimulate neurological cell proliferation and/or differentiation. Therefore, polynucleotides, polypeptides, agonists and/or antagonists of the invention may be used to treat and/or detect neurologic diseases. Moreover, polynucleotides or polypeptides, or agonists or antagonists of the invention, can be used as a marker or detector of a particular nervous system disease or disorder.

[0654] Examples of neurologic diseases which can be treated or detected with polynucleotides, polypeptides, agonists, and/or antagonists of the present invention include brain diseases, such as metabolic brain diseases which includes phenylketonuria such as maternal phenylketonuria, pyruvate carboxylase deficiency, pyruvate dehydrogenase complex deficiency, Wernicke's Encephalopathy, brain edema, brain neoplasms such as cerebellar neoplasms which include infratentorial neoplasms, cerebral ventricle neoplasms such as choroid plexus neoplasms, hypothalamic neoplasms, supratentorial neoplasms, canavan disease, cerebellar diseases such as cerebellar ataxia which include spinocerebellar degeneration such as ataxia telangiectasia, cerebellar dyssynergia, Friederich's Ataxia, Machado-Joseph Disease, olivopontocerebellar atrophy, cerebellar neoplasms such as infratentorial neoplasms, diffuse cerebral sclerosis such as encephalitis periaxialis, globoid cell leukodystrophy, metachromatic leukodystrophy and subacute sclerosing panencephalitis.

[0655] Additional neurologic diseases which can be treated or detected with polynucleotides, polypeptides, agonists, and/or antagonists of the present invention include cerebrovascular disorders (such as carotid artery diseases which include carotid artery thrombosis, carotid stenosis and Moyamoya Disease), cerebral amyloid angiopathy, cerebral aneurysm, cerebral anoxia, cerebral arteriosclerosis, cerebral arteriovenous

etc., using techniques known in the art and depending on the molecule to be measured; and motor neuron dysfunction may be measured by assessing the physical manifestation of motor neuron disorder, e.g., weakness, motor neuron conduction velocity, or functional disability.

[0650] In specific embodiments, motor neuron disorders that may be treated according to the invention include, but are not limited to, disorders such as infarction, infection, exposure to toxin, trauma, surgical damage, degenerative disease or malignancy that may affect motor neurons as well as other components of the nervous system, as well as disorders that selectively affect neurons such as amyotrophic lateral sclerosis, and including, but not limited to, progressive spinal muscular atrophy, progressive bulbar palsy, primary lateral sclerosis, infantile and juvenile muscular atrophy, progressive bulbar paralysis of childhood (Fazio-Londe syndrome), poliomyelitis and the post polio syndrome, and Hereditary Motorsensory Neuropathy (Charcot-Marie-Tooth Disease).

[0651] Further, polypeptides or polynucleotides of the invention may play a role in neuronal survival; synapse formation; conductance; neural differentiation, etc. Thus, compositions of the invention (including polynucleotides, polypeptides, and agonists or antagonists) may be used to diagnose and/or treat or prevent diseases or disorders associated with these roles, including, but not limited to, learning and/or cognition disorders. The compositions of the invention may also be useful in the treatment or prevention of neurodegenerative disease states and/or behavioural disorders. Such neurodegenerative disease states and/or behavioral disorders include, but are not limited to, Alzheimer's Disease, Parkinson's Disease, Huntington's Disease, Tourette Syndrome, schizophrenia, mania, dementia, paranoia, obsessive compulsive disorder, panic disorder, learning disabilities, ALS, psychoses, autism, and altered behaviors, including disorders in feeding, sleep patterns, balance, and perception. In addition, compositions of the invention may also play a role in the treatment, prevention and/or detection of developmental disorders associated with the developing embryo, or sexually-linked disorders.

[0652] Additionally, polypeptides, polynucleotides and/or agonists or antagonists of the invention, may be useful in protecting neural cells from diseases, damage, disorders, or injury, associated with cerebrovascular disorders including, but not limited to, carotid artery diseases (e.g., carotid artery thrombosis, carotid stenosis, or Moyamoya Disease), cerebral amyloid angiopathy, cerebral aneurysm, cerebral anoxia, cerebral arteriosclerosis,

exclusive aspect of this embodiment, the polypeptides, polynucleotides, or agonists or antagonists of the invention, are used to treat or prevent neural cell injury associated with cerebral ischemia. In another non-exclusive aspect of this embodiment, the polypeptides, polynucleotides, or agonists or antagonists of the invention are used to treat or prevent neural cell injury associated with cerebral infarction.

[0647] In another preferred embodiment, the polypeptides, polynucleotides, or agonists or antagonists of the invention are used to treat or prevent neural cell injury associated with a stroke. In a specific embodiment, the polypeptides, polynucleotides, or agonists or antagonists of the invention are used to treat or prevent cerebral neural cell injury associated with a stroke.

[0648] In another preferred embodiment, the polypeptides, polynucleotides, or agonists or antagonists of the invention are used to treat or prevent neural cell injury associated with a heart attack. In a specific embodiment, the polypeptides, polynucleotides, or agonists or antagonists of the invention are used to treat or prevent cerebral neural cell injury associated with a heart attack.

[0649] The compositions of the invention which are useful for treating or preventing a nervous system disorder may be selected by testing for biological activity in promoting the survival or differentiation of neurons. For example, and not by way of limitation, compositions of the invention which elicit any of the following effects may be useful according to the invention: (1) increased survival time of neurons in culture either in the presence or absence of hypoxia or hypoxic conditions; (2) increased sprouting of neurons in culture or *in vivo*; (3) increased production of a neuron-associated molecule in culture or *in vivo*, e.g., choline acetyltransferase or acetylcholinesterase with respect to motor neurons; or (4) decreased symptoms of neuron dysfunction *in vivo*. Such effects may be measured by any method known in the art. In preferred, non-limiting embodiments, increased survival of neurons may routinely be measured using a method set forth herein or otherwise known in the art, such as, for example, in Zhang *et al.*, *Proc Natl Acad Sci USA* 97:3637-42 (2000) or in Arakawa *et al.*, *J. Neurosci.*, 10:3507-15 (1990); increased sprouting of neurons may be detected by methods known in the art, such as, for example, the methods set forth in Pestronk *et al.*, *Exp. Neurol.*, 70:65-82 (1980), or Brown *et al.*, *Ann. Rev. Neurosci.*, 4:17-42 (1981); increased production of neuron-associated molecules may be measured by bioassay, enzymatic assay, antibody binding, Northern blot assay,

(including spinal cord, brain) or peripheral nervous systems: (1) ischemic lesions, in which a lack of oxygen in a portion of the nervous system results in neuronal injury or death, including cerebral infarction or ischemia, or spinal cord infarction or ischemia; (2) traumatic lesions, including lesions caused by physical injury or associated with surgery, for example, lesions which sever a portion of the nervous system, or compression injuries; (3) malignant lesions, in which a portion of the nervous system is destroyed or injured by malignant tissue which is either a nervous system associated malignancy or a malignancy derived from non-nervous system tissue; (4) infectious lesions, in which a portion of the nervous system is destroyed or injured as a result of infection, for example, by an abscess or associated with infection by human immunodeficiency virus, herpes zoster, or herpes simplex virus or with Lyme disease, tuberculosis, or syphilis; (5) degenerative lesions, in which a portion of the nervous system is destroyed or injured as a result of a degenerative process including but not limited to, degeneration associated with Parkinson's disease, Alzheimer's disease, Huntington's chorea, or amyotrophic lateral sclerosis (ALS); (6) lesions associated with nutritional diseases or disorders, in which a portion of the nervous system is destroyed or injured by a nutritional disorder or disorder of metabolism including, but not limited to, vitamin B12 deficiency, folic acid deficiency, Wernicke disease, tobacco-alcohol amblyopia, Marchiafava-Bignami disease (primary degeneration of the corpus callosum), and alcoholic cerebellar degeneration; (7) neurological lesions associated with systemic diseases including, but not limited to, diabetes (diabetic neuropathy, Bell's palsy), systemic lupus erythematosus, carcinoma, or sarcoidosis; (8) lesions caused by toxic substances including alcohol, lead, or particular neurotoxins; and (9) demyelinated lesions in which a portion of the nervous system is destroyed or injured by a demyelinating disease including, but not limited to, multiple sclerosis, human immunodeficiency virus-associated myelopathy, transverse myelopathy or various etiologies, progressive multifocal leukoencephalopathy, and central pontine myelinolysis.

[0646] In one embodiment, the polypeptides, polynucleotides, or agonists or antagonists of the invention are used to protect neural cells from the damaging effects of hypoxia. In a further preferred embodiment, the polypeptides, polynucleotides, or agonists or antagonists of the invention are used to protect neural cells from the damaging effects of cerebral hypoxia. According to this embodiment, the compositions of the invention are used to treat or prevent neural cell injury associated with cerebral hypoxia. In one non-

burns, that cause necrosis of the bronchiolar epithelium and alveoli could be effectively treated using polynucleotides or polypeptides, agonists or antagonists of the present invention. Also, polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could be used to stimulate the proliferation of and differentiation of type II pneumocytes, which may help treat or prevent disease such as hyaline membrane diseases, such as infant respiratory distress syndrome and bronchopulmonary dysplasia, in premature infants.

[0643] Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could stimulate the proliferation and differentiation of hepatocytes and, thus, could be used to alleviate or treat liver diseases and pathologies such as fulminant liver failure caused by cirrhosis, liver damage caused by viral hepatitis and toxic substances (i.e., acetaminophen, carbon tetrachloride and other hepatotoxins known in the art).

[0644] In addition, polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could be used to treat or prevent the onset of diabetes mellitus. In patients with newly diagnosed Types I and II diabetes, where some islet cell function remains, polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could be used to maintain the islet function so as to alleviate, delay or prevent permanent manifestation of the disease. Also, polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could be used as an auxiliary in islet cell transplantation to improve or promote islet cell function.

Neural Activity and Neurological Diseases

[0645] The polynucleotides, polypeptides and agonists or antagonists of the invention may be used for the diagnosis and/or treatment of diseases, disorders, damage or injury of the brain and/or nervous system. Nervous system disorders that can be treated with the compositions of the invention (e.g., polypeptides, polynucleotides, and/or agonists or antagonists), include, but are not limited to, nervous system injuries, and diseases or disorders which result in either a disconnection of axons, a diminution or degeneration of neurons, or demyelination. Nervous system lesions which may be treated in a patient (including human and non-human mammalian patients) according to the methods of the invention, include but are not limited to, the following lesions of either the central

cytoprotective effect on the small intestine mucosa. Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, may also stimulate healing of mucositis (mouth ulcers) that result from chemotherapy and viral infections.

[0641] Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could further be used in full regeneration of skin in full and partial thickness skin defects, including burns, (i.e., repopulation of hair follicles, sweat glands, and sebaceous glands), treatment of other skin defects such as psoriasis. Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could be used to treat epidermolysis bullosa, a defect in adherence of the epidermis to the underlying dermis which results in frequent, open and painful blisters by accelerating reepithelialization of these lesions. Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could also be used to treat gastric and duodenal ulcers and help heal by scar formation of the mucosal lining and regeneration of glandular mucosa and duodenal mucosal lining more rapidly. Inflammatory bowel diseases, such as Crohn's disease and ulcerative colitis, are diseases which result in destruction of the mucosal surface of the small or large intestine, respectively. Thus, polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could be used to promote the resurfacing of the mucosal surface to aid more rapid healing and to prevent progression of inflammatory bowel disease. Treatment with polynucleotides or polypeptides, agonists or antagonists of the present invention, is expected to have a significant effect on the production of mucus throughout the gastrointestinal tract and could be used to protect the intestinal mucosa from injurious substances that are ingested or following surgery. Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could be used to treat diseases associated with the under expression.

[0642] Moreover, polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could be used to prevent and heal damage to the lungs due to various pathological states. Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, which could stimulate proliferation and differentiation and promote the repair of alveoli and bronchiolar epithelium to prevent or treat acute or chronic lung damage. For example, emphysema, which results in the progressive loss of alveoli, and inhalation injuries, i.e., resulting from smoke inhalation and

venous stasis ulcers, burns resulting from heat exposure or chemicals, and other abnormal wound healing conditions such as uremia, malnutrition, vitamin deficiencies and complications associated with systemic treatment with steroids, radiation therapy and antineoplastic drugs and antimetabolites. Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could be used to promote dermal reestablishment subsequent to dermal loss

[0638] Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could be used to increase the adherence of skin grafts to a wound bed and to stimulate re-epithelialization from the wound bed. The following are types of grafts that polynucleotides or polypeptides, agonists or antagonists of the present invention, could be used to increase adherence to a wound bed: autografts, artificial skin, allografts, autodermic graft, autoepidermic grafts, avascular grafts, Blair-Brown grafts, bone graft, brephoplastic grafts, cutis graft, delayed graft, dermic graft, epidermic graft, fascia graft, full thickness graft, heterologous graft, xenograft, homologous graft, hyperplastic graft, lamellar graft, mesh graft, mucosal graft, Ollier-Thiersch graft, omentopial graft, patch graft, pedicle graft, penetrating graft, split skin graft, thick split graft. Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, can be used to promote skin strength and to improve the appearance of aged skin.

[0639] It is believed that polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, will also produce changes in hepatocyte proliferation, and epithelial cell proliferation in the lung, breast, pancreas, stomach, small intestine, and large intestine. Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could promote proliferation of epithelial cells such as sebocytes, hair follicles, hepatocytes, type II pneumocytes, mucin-producing goblet cells, and other epithelial cells and their progenitors contained within the skin, lung, liver, and gastrointestinal tract. Polynucleotides or polypeptides, agonists or antagonists of the present invention, may promote proliferation of endothelial cells, keratinocytes, and basal keratinocytes.

[0640] Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, could also be used to reduce the side effects of gut toxicity that result from radiation, chemotherapy treatments or viral infections. Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, may have a

cell carcinoma, basal cell carcinoma, adenocarcinoma, sweat gland carcinoma, sebaceous gland carcinoma, papillary carcinoma, papillary adenocarcinomas, cystadenocarcinoma, medullary carcinoma, bronchogenic carcinoma, renal cell carcinoma, hepatoma, bile duct carcinoma, choriocarcinoma, seminoma, embryonal carcinoma, Wilm's tumor, cervical cancer, testicular tumor, lung carcinoma, small cell lung carcinoma, bladder carcinoma, epithelial carcinoma, glioma, astrocytoma, medulloblastoma, craniopharyngioma, ependymoma, pinealoma, hemangioblastoma, acoustic neuroma, oligodendroglioma, meningioma, melanoma, neuroblastoma, and retinoblastoma.

[0636] Diseases associated with increased apoptosis that could be treated, prevented, diagnosed, and/or prognosed using polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, include, but are not limited to, AIDS; neurodegenerative disorders (such as Alzheimer's disease, Parkinson's disease, Amyotrophic lateral sclerosis, Retinitis pigmentosa, Cerebellar degeneration and brain tumor or prior associated disease); autoimmune disorders (such as, multiple sclerosis, Sjogren's syndrome, Hashimoto's thyroiditis, biliary cirrhosis, Behcet's disease, Crohn's disease, polymyositis, systemic lupus erythematosus and immune-related glomerulonephritis and rheumatoid arthritis) myelodysplastic syndromes (such as aplastic anemia), graft v. host disease, ischemic injury (such as that caused by myocardial infarction, stroke and reperfusion injury), liver injury (e.g., hepatitis related liver injury, ischemia/reperfusion injury, cholestasis (bile duct injury) and liver cancer); toxin-induced liver disease (such as that caused by alcohol), septic shock, cachexia and anorexia.

Wound Healing and Epithelial Cell Proliferation

[0637] In accordance with yet a further aspect of the present invention, there is provided a process for utilizing polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, for therapeutic purposes, for example, to stimulate epithelial cell proliferation and basal keratinocytes for the purpose of wound healing, and to stimulate hair follicle production and healing of dermal wounds. Polynucleotides or polypeptides, as well as agonists or antagonists of the present invention, may be clinically useful in stimulating wound healing including surgical wounds, excisional wounds, deep wounds involving damage of the dermis and epidermis, eye tissue wounds, dental tissue wounds, oral cavity wounds, diabetic ulcers, dermal ulcers, cubitus ulcers, arterial ulcers,

Diseases at the Cellular Level

[0633] Diseases associated with increased cell survival or the inhibition of apoptosis that could be treated, prevented, diagnosed, and/or prognosed using polynucleotides or polypeptides, as well as antagonists or agonists of the present invention, include cancers (such as follicular lymphomas, carcinomas with p53 mutations, and hormone-dependent tumors, including, but not limited to colon cancer, cardiac tumors, pancreatic cancer, melanoma, retinoblastoma, glioblastoma, lung cancer, intestinal cancer, testicular cancer, stomach cancer, neuroblastoma, myxoma, myoma, lymphoma, endothelioma, osteoblastoma, osteoclastoma, osteosarcoma, chondrosarcoma, adenoma, breast cancer, prostate cancer, Kaposi's sarcoma and ovarian cancer); autoimmune disorders (such as, multiple sclerosis, Sjogren's syndrome, Hashimoto's thyroiditis, biliary cirrhosis, Behcet's disease, Crohn's disease, polymyositis, systemic lupus erythematosus and immune-related glomerulonephritis and rheumatoid arthritis) and viral infections (such as herpes viruses, pox viruses and adenoviruses), inflammation, graft v. host disease, acute graft rejection, and chronic graft rejection.

[0634] In preferred embodiments, polynucleotides, polypeptides, and/or antagonists of the invention are used to inhibit growth, progression, and/or metasis of cancers, in particular those listed above.

[0635] Additional diseases or conditions associated with increased cell survival that could be treated or detected by polynucleotides or polypeptides, or agonists or antagonists of the present invention include, but are not limited to, progression, and/or metastases of malignancies and related disorders such as leukemia (including acute leukemias (e.g., acute lymphocytic leukemia, acute myelocytic leukemia (including myeloblastic, promyelocytic, myelomonocytic, monocytic, and erythroleukemia)) and chronic leukemias (e.g., chronic myelocytic (granulocytic) leukemia and chronic lymphocytic leukemia)), polycythemia vera, lymphomas (e.g., Hodgkin's disease and non-Hodgkin's disease), multiple myeloma, Waldenstrom's macroglobulinemia, heavy chain disease, and solid tumors including, but not limited to, sarcomas and carcinomas such as fibrosarcoma, myxosarcoma, liposarcoma, chondrosarcoma, osteogenic sarcoma, chordoma, angiosarcoma, endotheliosarcoma, lymphangiosarcoma, lymphangioendotheliosarcoma, synovioma, mesothelioma, Ewing's tumor, leiomyosarcoma, rhabdomyosarcoma, colon carcinoma, pancreatic cancer, breast cancer, ovarian cancer, prostate cancer, squamous

[0631] Representative examples of tungsten and molybdenum complexes also include oxo complexes. Suitable oxo tungsten complexes include tungstate and tungsten oxide complexes. Suitable tungstate complexes include ammonium tungstate, calcium tungstate, sodium tungstate dihydrate, and tungstic acid. Suitable tungsten oxides include tungsten (IV) oxide and tungsten (VI) oxide. Suitable oxo molybdenum complexes include molybdate, molybdenum oxide, and molybdenyl complexes. Suitable molybdate complexes include ammonium molybdate and its hydrates, sodium molybdate and its hydrates, and potassium molybdate and its hydrates. Suitable molybdenum oxides include molybdenum (VI) oxide, molybdenum (VI) oxide, and molybdic acid. Suitable molybdenyl complexes include, for example, molybdenyl acetylacetonate. Other suitable tungsten and molybdenum complexes include hydroxo derivatives derived from, for example, glycerol, tartaric acid, and sugars.

[0632] A wide variety of other anti-angiogenic factors may also be utilized within the context of the present invention. Representative examples include platelet factor 4; protamine sulphate; sulphated chitin derivatives (prepared from queen crab shells), (Murata et al., Cancer Res. 51:22-26, 1991); Sulphated Polysaccharide Peptidoglycan Complex (SP- PG) (the function of this compound may be enhanced by the presence of steroids such as estrogen, and tamoxifen citrate); Staurosporine; modulators of matrix metabolism, including for example, proline analogs, cishydroxyproline, d,L-3,4-dehydroproline, Thiaproline, alpha,alpha-dipyridyl, aminopropionitrile fumarate; 4-propyl-5-(4-pyridinyl)-2(3H)-oxazolone; Methotrexate; Mitoxantrone; Heparin; Interferons; 2 Macroglobulin-serum; ChIMP-3 (Pavloff et al., J. Bio. Chem. 267:17321-17326, 1992); Chymostatin (Tomkinson et al., Biochem J. 286:475-480, 1992); Cyclodextrin Tetradecasulfate; Eponemycin; Camptothecin; Fumagillin (Ingber et al., Nature 348:555-557, 1990); Gold Sodium Thiomalate ("GST"; Matsubara and Ziff, J. Clin. Invest. 79:1440-1446, 1987); anticollagenase-serum; alpha2-antiplasmin (Holmes et al., J. Biol. Chem. 262(4):1659-1664, 1987); Bisantrene (National Cancer Institute); Lobenzarit disodium (N-(2)-carboxyphenyl-4- chloroanthronilic acid disodium or "CCA"; Takeuchi et al., Agents Actions 36:312-316, 1992); Thalidomide; Angostatic steroid; AGM-1470; carboxynaminolmidazole; and metalloproteinase inhibitors such as BB94.

agonist and/or agonist to the resection margins of a tumor subsequent to excision, such that the local recurrence of cancer and the formation of new blood vessels at the site is inhibited. Within one embodiment of the invention, the anti-angiogenic compound is administered directly to the tumor excision site (e.g., applied by swabbing, brushing or otherwise coating the resection margins of the tumor with the anti-angiogenic compound). Alternatively, the anti-angiogenic compounds may be incorporated into known surgical pastes prior to administration. Within particularly preferred embodiments of the invention, the anti-angiogenic compounds are applied after hepatic resections for malignancy, and after neurosurgical operations.

[0627] Within one aspect of the present invention, polynucleotides, polypeptides, agonists and/or agonists may be administered to the resection margin of a wide variety of tumors, including for example, breast, colon, brain and hepatic tumors. For example, within one embodiment of the invention, anti-angiogenic compounds may be administered to the site of a neurological tumor subsequent to excision, such that the formation of new blood vessels at the site are inhibited.

[0628] The polynucleotides, polypeptides, agonists and/or agonists of the present invention may also be administered along with other anti-angiogenic factors. Representative examples of other anti-angiogenic factors include: Anti-Invasive Factor, retinoic acid and derivatives thereof, paclitaxel, Suramin, Tissue Inhibitor of Metalloproteinase-1, Tissue Inhibitor of Metalloproteinase-2, Plasminogen Activator Inhibitor-1, Plasminogen Activator Inhibitor-2, and various forms of the lighter "d group" transition metals.

[0629] Lighter "d group" transition metals include, for example, vanadium, molybdenum, tungsten, titanium, niobium, and tantalum species. Such transition metal species may form transition metal complexes. Suitable complexes of the above-mentioned transition metal species include oxo transition metal complexes.

[0630] Representative examples of vanadium complexes include oxo vanadium complexes such as vanadate and vanadyl complexes. Suitable vanadate complexes include metavanadate and orthovanadate complexes such as, for example, ammonium metavanadate, sodium metavanadate, and sodium orthovanadate. Suitable vanadyl complexes include, for example, vanadyl acetylacetonate and vanadyl sulfate including vanadyl sulfate hydrates such as vanadyl sulfate mono- and trihydrates.

(keloids), nonunion fractures, scleroderma, trachoma, vascular adhesions, myocardial angiogenesis, coronary collaterals, cerebral collaterals, arteriovenous malformations, ischemic limb angiogenesis, Osler-Webber Syndrome, plaque neovascularization, telangiectasia, hemophiliac joints, angiofibroma fibromuscular dysplasia, wound granulation, Crohn's disease, atherosclerosis, birth control agent by preventing vascularization required for embryo implantation controlling menstruation, diseases that have angiogenesis as a pathologic consequence such as cat scratch disease (*Rochelie minalia quintosa*), ulcers (*Helicobacter pylori*), Bartonellosis and bacillary angiomatosis.

[0623] In one aspect of the birth control method, an amount of the compound sufficient to block embryo implantation is administered before or after intercourse and fertilization have occurred, thus providing an effective method of birth control, possibly a "morning after" method. Polynucleotides, polypeptides, agonists and/or agonists may also be used in controlling menstruation or administered as either a peritoneal lavage fluid or for peritoneal implantation in the treatment of endometriosis.

[0624] Polynucleotides, polypeptides, agonists and/or agonists of the present invention may be incorporated into surgical sutures in order to prevent stitch granulomas.

[0625] Polynucleotides, polypeptides, agonists and/or agonists may be utilized in a wide variety of surgical procedures. For example, within one aspect of the present invention a compositions (in the form of, for example, a spray or film) may be utilized to coat or spray an area prior to removal of a tumor, in order to isolate normal surrounding tissues from malignant tissue, and/or to prevent the spread of disease to surrounding tissues. Within other aspects of the present invention, compositions (e.g., in the form of a spray) may be delivered via endoscopic procedures in order to coat tumors, or inhibit angiogenesis in a desired locale. Within yet other aspects of the present invention, surgical meshes which have been coated with anti- angiogenic compositions of the present invention may be utilized in any procedure wherein a surgical mesh might be utilized. For example, within one embodiment of the invention a surgical mesh laden with an anti-angiogenic composition may be utilized during abdominal cancer resection surgery (e.g., subsequent to colon resection) in order to provide support to the structure, and to release an amount of the anti-angiogenic factor.

[0626] Within further aspects of the present invention, methods are provided for treating tumor excision sites, comprising administering a polynucleotide, polypeptide,

forms of neovascular glaucoma. Within other embodiments, the compound may be implanted by injection into the region of the anterior chamber angle. Within other embodiments, the compound may also be placed in any location such that the compound is continuously released into the aqueous humor. Within another aspect of the present invention, methods are provided for treating proliferative diabetic retinopathy, comprising the step of administering to a patient a therapeutically effective amount of a polynucleotide, polypeptide, antagonist and/or agonist to the eyes, such that the formation of blood vessels is inhibited.

[0619] Within particularly preferred embodiments of the invention, proliferative diabetic retinopathy may be treated by injection into the aqueous humor or the vitreous, in order to increase the local concentration of the polynucleotide, polypeptide, antagonist and/or agonist in the retina. Preferably, this treatment should be initiated prior to the acquisition of severe disease requiring photocoagulation.

[0620] Within another aspect of the present invention, methods are provided for treating retrolental fibroplasia, comprising the step of administering to a patient a therapeutically effective amount of a polynucleotide, polypeptide, antagonist and/or agonist to the eye, such that the formation of blood vessels is inhibited. The compound may be administered topically, via intravitreal injection and/or via intraocular implants.

[0621] Additionally, disorders which can be treated with the polynucleotides, polypeptides, agonists and/or antagonists include, but are not limited to, hemangioma, arthritis, psoriasis, angiofibroma, atherosclerotic plaques, delayed wound healing, granulations, hemophilic joints, hypertrophic scars, nonunion fractures, Osler-Weber syndrome, pyogenic granuloma, scleroderma, trachoma, and vascular adhesions.

[0622] Moreover, disorders and/or states, which can be treated, prevented, diagnosed, and/or prognosed with the polynucleotides, polypeptides, agonists and/or antagonists of the invention include, but are not limited to, solid tumors, blood born tumors such as leukemias, tumor metastasis, Kaposi's sarcoma, benign tumors, for example hemangiomas, acoustic neuromas, neurofibromas, trachomas, and pyogenic granulomas, rheumatoid arthritis, psoriasis, ocular angiogenic diseases, for example, diabetic retinopathy, retinopathy of prematurity, macular degeneration, corneal graft rejection, neovascular glaucoma, retrolental fibroplasia, rubeosis, retinoblastoma, and uveitis, delayed wound healing, endometriosis, vasculogenesis, granulations, hypertrophic scars

deficiency states, and as a complication of wearing contact lenses.

[0616] Within particularly preferred embodiments of the invention, may be prepared for topical administration in saline (combined with any of the preservatives and antimicrobial agents commonly used in ocular preparations), and administered in eyedrop form. The solution or suspension may be prepared in its pure form and administered several times daily. Alternatively, anti-angiogenic compositions, prepared as described above, may also be administered directly to the cornea. Within preferred embodiments, the anti-angiogenic composition is prepared with a muco-adhesive polymer which binds to cornea. Within further embodiments, the anti-angiogenic factors or anti-angiogenic compositions may be utilized as an adjunct to conventional steroid therapy. Topical therapy may also be useful prophylactically in corneal lesions which are known to have a high probability of inducing an angiogenic response (such as chemical burns). In these instances the treatment, likely in combination with steroids, may be instituted immediately to help prevent subsequent complications.

[0617] Within other embodiments, the compounds described above may be injected directly into the corneal stroma by an ophthalmologist under microscopic guidance. The preferred site of injection may vary with the morphology of the individual lesion, but the goal of the administration would be to place the composition at the advancing front of the vasculature (i.e., interspersed between the blood vessels and the normal cornea). In most cases this would involve perilimbal corneal injection to "protect" the cornea from the advancing blood vessels. This method may also be utilized shortly after a corneal insult in order to prophylactically prevent corneal neovascularization. In this situation the material could be injected in the perilimbal cornea interspersed between the corneal lesion and its undesired potential limbal blood supply. Such methods may also be utilized in a similar fashion to prevent capillary invasion of transplanted corneas. In a sustained-release form injections might only be required 2-3 times per year. A steroid could also be added to the injection solution to reduce inflammation resulting from the injection itself.

[0618] Within another aspect of the present invention, methods are provided for treating neovascular glaucoma, comprising the step of administering to a patient a therapeutically effective amount of a polynucleotide, polypeptide, antagonist and/or agonist to the eye, such that the formation of blood vessels is inhibited. In one embodiment, the compound may be administered topically to the eye in order to treat early

scar or keloid.

[0613] Within one embodiment of the present invention polynucleotides, polypeptides, antagonists and/or agonists of the invention are directly injected into a hypertrophic scar or keloid, in order to prevent the progression of these lesions. This therapy is of particular value in the prophylactic treatment of conditions which are known to result in the development of hypertrophic scars and keloids (e.g., burns), and is preferably initiated after the proliferative phase has had time to progress (approximately 14 days after the initial injury), but before hypertrophic scar or keloid development. As noted above, the present invention also provides methods for treating neovascular diseases of the eye, including for example, corneal neovascularization, neovascular glaucoma, proliferative diabetic retinopathy, retrolental fibroplasia and macular degeneration.

[0614] Moreover, Ocular disorders associated with neovascularization which can be treated with the polynucleotides and polypeptides of the present invention (including agonists and/or antagonists) include, but are not limited to: neovascular glaucoma, diabetic retinopathy, retinoblastoma, retrolental fibroplasia, uveitis, retinopathy of prematurity macular degeneration, corneal graft neovascularization, as well as other eye inflammatory diseases, ocular tumors and diseases associated with choroidal or iris neovascularization. See, e.g., reviews by Waltman *et al.*, *Am. J. Ophthalmol.* 85:704-710 (1978) and Gartner *et al.*, *Surv. Ophthalmol.* 22:291-312 (1978).

[0615] Thus, within one aspect of the present invention methods are provided for treating neovascular diseases of the eye such as corneal neovascularization (including corneal graft neovascularization), comprising the step of administering to a patient a therapeutically effective amount of a compound (as described above) to the cornea, such that the formation of blood vessels is inhibited. Briefly, the cornea is a tissue which normally lacks blood vessels. In certain pathological conditions however, capillaries may extend into the cornea from the pericorneal vascular plexus of the limbus. When the cornea becomes vascularized, it also becomes clouded, resulting in a decline in the patient's visual acuity. Visual loss may become complete if the cornea completely opacitates. A wide variety of disorders can result in corneal neovascularization, including for example, corneal infections (e.g., trachoma, herpes simplex keratitis, leishmaniasis and onchocerciasis), immunological processes (e.g., graft rejection and Stevens-Johnson's syndrome), alkali burns, trauma, inflammation (of any cause), toxic and nutritional

antagonists and/or agonists include, but are not limited to solid tumors, including prostate, lung, breast, ovarian, stomach, pancreas, larynx, esophagus, testes, liver, parotid, biliary tract, colon, rectum, cervix, uterus, endometrium, kidney, bladder, thyroid cancer; primary tumors and metastases; melanomas; glioblastoma; Kaposi's sarcoma; leiomyosarcoma; non-small cell lung cancer; colorectal cancer; advanced malignancies; and blood born tumors such as leukemias. For example, polynucleotides, polypeptides, antagonists and/or agonists may be delivered topically, in order to treat cancers such as skin cancer, head and neck tumors, breast tumors, and Kaposi's sarcoma.

[0610] Within yet other aspects, polynucleotides, polypeptides, antagonists and/or agonists may be utilized to treat superficial forms of bladder cancer by, for example, intravesical administration. Polynucleotides, polypeptides, antagonists and/or agonists may be delivered directly into the tumor, or near the tumor site, via injection or a catheter. Of course, as the artisan of ordinary skill will appreciate, the appropriate mode of administration will vary according to the cancer to be treated. Other modes of delivery are discussed herein.

[0611] Polynucleotides, polypeptides, antagonists and/or agonists may be useful in treating other disorders, besides cancers, which involve angiogenesis. These disorders include, but are not limited to: benign tumors, for example hemangiomas, acoustic neuromas, neurofibromas, trachomas, and pyogenic granulomas; arteriosclerotic plaques; ocular angiogenic diseases, for example, diabetic retinopathy, retinopathy of prematurity, macular degeneration, corneal graft rejection, neovascular glaucoma, retrolental fibroplasia, rubeosis, retinoblastoma, uveitis and Pterygia (abnormal blood vessel growth) of the eye; rheumatoid arthritis; psoriasis; delayed wound healing; endometriosis; vasculogenesis; granulations; hypertrophic scars (keloids); nonunion fractures; scleroderma; trachoma; vascular adhesions; myocardial angiogenesis; coronary collaterals; cerebral collaterals; arteriovenous malformations; ischemic limb angiogenesis; Osler-Webber Syndrome; plaque neovascularization; telangiectasia; hemophilic joints; angiofibroma; fibromuscular dysplasia; wound granulation; Crohn's disease; and atherosclerosis.

[0612] For example, within one aspect of the present invention methods are provided for treating hypertrophic scars and keloids, comprising the step of administering a polynucleotide, polypeptide, antagonist and/or agonist of the invention to a hypertrophic

Anti-Angiogenesis Activity

[0608] The naturally occurring balance between endogenous stimulators and inhibitors of angiogenesis is one in which inhibitory influences predominate. Rastinejad *et al.*, *Cell* 56:345-355 (1989). In those rare instances in which neovascularization occurs under normal physiological conditions, such as wound healing, organ regeneration, embryonic development, and female reproductive processes, angiogenesis is stringently regulated and spatially and temporally delimited. Under conditions of pathological angiogenesis such as that characterizing solid tumor growth, these regulatory controls fail. Unregulated angiogenesis becomes pathologic and sustains progression of many neoplastic and non-neoplastic diseases. A number of serious diseases are dominated by abnormal neovascularization including solid tumor growth and metastases, arthritis, some types of eye disorders, and psoriasis. See, e.g., reviews by Moses *et al.*, *Biotech.* 9:630-634 (1991); Folkman *et al.*, *N. Engl. J. Med.*, 333:1757-1763 (1995); Auerbach *et al.*, *J. Microvasc. Res.* 29:401-411 (1985); Folkman, *Advances in Cancer Research*, eds. Klein and Weinhouse, Academic Press, New York, pp. 175-203 (1985); Patz, *Am. J. Ophthalmol.* 94:715-743 (1982); and Folkman *et al.*, *Science* 221:719-725 (1983). In a number of pathological conditions, the process of angiogenesis contributes to the disease state. For example, significant data have accumulated which suggest that the growth of solid tumors is dependent on angiogenesis. Folkman and Klagsbrun, *Science* 235:442-447 (1987).

[0609] The present invention provides for treatment of diseases or disorders associated with neovascularization by administration of the polynucleotides and/or polypeptides of the invention, as well as agonists or antagonists of the present invention. Malignant and metastatic conditions which can be treated with the polynucleotides and polypeptides, or agonists or antagonists of the invention include, but are not limited to, malignancies, solid tumors, and cancers described herein and otherwise known in the art (for a review of such disorders, see Fishman *et al.*, *Medicine*, 2d Ed., J. B. Lippincott Co., Philadelphia (1985)). Thus, the present invention provides a method of treating an angiogenesis-related disease and/or disorder, comprising administering to an individual in need thereof a therapeutically effective amount of a polynucleotide, polypeptide, antagonist and/or agonist of the invention. For example, polynucleotides, polypeptides, antagonists and/or agonists may be utilized in a variety of additional methods in order to therapeutically treat a cancer or tumor. Cancers which may be treated with polynucleotides, polypeptides,

bacterial pneumonia (e.g., *Streptococcus pneumoniae* (pneumococcal pneumonia), *Staphylococcus aureus* (staphylococcal pneumonia), Gram-negative bacterial pneumonia (caused by, e.g., *Klebsiella* and *Pseudomonas spp.*), *Mycoplasma pneumoniae* pneumonia, *Hemophilus influenzae* pneumonia, *Legionella pneumophila* (Legionnaires' disease), and *Chlamydia psittaci* (Psittacosis)), and viral pneumonia (e.g., influenza, chickenpox (varicella).

[0607] Additional diseases and disorders of the respiratory system include, but are not limited to bronchiolitis, polio (poliomyelitis), croup, respiratory syncytial viral infection, mumps, erythema infectiosum (fifth disease), roseola infantum, progressive rubella panencephalitis, german measles, and subacute sclerosing panencephalitis), fungal pneumonia (e.g., Histoplasmosis, Coccidioidomycosis, Blastomycosis, fungal infections in people with severely suppressed immune systems (e.g., cryptococcosis, caused by *Cryptococcus neoformans*; aspergillosis, caused by *Aspergillus spp.*; candidiasis, caused by *Candida*; and mucormycosis)), *Pneumocystis carinii* (pneumocystis pneumonia), atypical pneumonias (e.g., *Mycoplasma* and *Chlamydia spp.*), opportunistic infection pneumonia, nosocomial pneumonia, chemical pneumonitis, and aspiration pneumonia, pleural disorders (e.g., pleurisy, pleural effusion, and pneumothorax (e.g., simple spontaneous pneumothorax, complicated spontaneous pneumothorax, tension pneumothorax)), obstructive airway diseases (e.g., asthma, chronic obstructive pulmonary disease (COPD), emphysema, chronic or acute bronchitis), occupational lung diseases (e.g., silicosis, black lung (coal workers' pneumoconiosis), asbestosis, berylliosis, occupational asthma, byssinosis, and benign pneumoconioses), Infiltrative Lung Disease (e.g., pulmonary fibrosis (e.g., fibrosing alveolitis, usual interstitial pneumonia), idiopathic pulmonary fibrosis, desquamative interstitial pneumonia, lymphoid interstitial pneumonia, histiocytosis X (e.g., Letterer-Siwe disease, Hand-Schüller-Christian disease, eosinophilic granuloma), idiopathic pulmonary hemosiderosis, sarcoidosis and pulmonary alveolar proteinosis), Acute respiratory distress syndrome (also called, e.g., adult respiratory distress syndrome), edema, pulmonary embolism, bronchitis (e.g., viral, bacterial), bronchiectasis, atelectasis, lung abscess (caused by, e.g., *Staphylococcus aureus* or *Legionella pneumophila*), and cystic fibrosis.

[0603] Ischemic disorders include, but are not limited to, cerebral ischemia, ischemic colitis, compartment syndromes, anterior compartment syndrome, myocardial ischemia, reperfusion injuries, and peripheral limb ischemia. Vasculitis includes, but is not limited to, aortitis, arteritis, Behcet's Syndrome, Churg-Strauss Syndrome, mucocutaneous lymph node syndrome, thromboangiitis obliterans, hypersensitivity vasculitis, Schoenlein-Henoch purpura, allergic cutaneous vasculitis, and Wegener's granulomatosis.

[0604] Polypeptides may be administered using any method known in the art, including, but not limited to, direct needle injection at the delivery site, intravenous injection, topical administration, catheter infusion, biolistic injectors, particle accelerators, gelfoam sponge depots, other commercially available depot materials, osmotic pumps, oral or suppository solid pharmaceutical formulations, decanting or topical applications during surgery, aerosol delivery. Such methods are known in the art. Polypeptides may be administered as part of a Therapeutic, described in more detail below. Methods of delivering polynucleotides are described in more detail herein.

Respiratory Disorders

[0605] Polynucleotides or polypeptides, or agonists or antagonists of the present invention may be used to treat, prevent, diagnose, and/or prognose diseases and/or disorders of the respiratory system.

[0606] Diseases and disorders of the respiratory system include, but are not limited to, nasal vestibulitis, nonallergic rhinitis (e.g., acute rhinitis, chronic rhinitis, atrophic rhinitis, vasomotor rhinitis), nasal polyps, and sinusitis, juvenile angiofibromas, cancer of the nose and juvenile papillomas, vocal cord polyps, nodules (singer's nodules), contact ulcers, vocal cord paralysis, laryngoceles, pharyngitis (e.g., viral and bacterial), tonsillitis, tonsillar cellulitis, parapharyngeal abscess, laryngitis, laryngoceles, and throat cancers (e.g., cancer of the nasopharynx, tonsil cancer, larynx cancer), lung cancer (e.g., squamous cell carcinoma, small cell (oat cell) carcinoma, large cell carcinoma, and adenocarcinoma), allergic disorders (eosinophilic pneumonia, hypersensitivity pneumonitis (e.g., extrinsic allergic alveolitis, allergic interstitial pneumonitis, organic dust pneumoconiosis, allergic bronchopulmonary aspergillosis, asthma, Wegener's granulomatosis (granulomatous vasculitis), Goodpasture's syndrome)), pneumonia (e.g.,

[0598] Cardiovascular diseases also include vascular diseases such as aneurysms, angiodysplasia, angiomas, bacillary angiomas, Hippel-Lindau Disease, Klippel-Trenaunay-Weber Syndrome, Sturge-Weber Syndrome, angioneurotic edema, aortic diseases, Takayasu's Arteritis, aortitis, Leriche's Syndrome, arterial occlusive diseases, arteritis, enarteritis, polyarteritis nodosa, cerebrovascular disorders, diabetic angiopathies, diabetic retinopathy, embolisms, thrombosis, erythromelalgia, hemorrhoids, hepatic veno-occlusive disease, hypertension, hypotension, ischemia, peripheral vascular diseases, phlebitis, pulmonary veno-occlusive disease, Raynaud's disease, CREST syndrome, retinal vein occlusion, Scimitar syndrome, superior vena cava syndrome, telangiectasia, ataxia telangiectasia, hereditary hemorrhagic telangiectasia, varicocele, varicose veins, varicose ulcer, vasculitis, and venous insufficiency.

[0599] Aneurysms include, but are not limited to, dissecting aneurysms, false aneurysms, infected aneurysms, ruptured aneurysms, aortic aneurysms, cerebral aneurysms, coronary aneurysms, heart aneurysms, and iliac aneurysms.

[0600] Arterial occlusive diseases include, but are not limited to, arteriosclerosis, intermittent claudication, carotid stenosis, fibromuscular dysplasias, mesenteric vascular occlusion, Moyamoya disease, renal artery obstruction, retinal artery occlusion, and thromboangiitis obliterans.

[0601] Cerebrovascular disorders include, but are not limited to, carotid artery diseases, cerebral amyloid angiopathy, cerebral aneurysm, cerebral anoxia, cerebral arteriosclerosis, cerebral arteriovenous malformation, cerebral artery diseases, cerebral embolism and thrombosis, carotid artery thrombosis, sinus thrombosis, Wallenberg's syndrome, cerebral hemorrhage, epidural hematoma, subdural hematoma, subarachnoid hemorrhage, cerebral infarction, cerebral ischemia (including transient), subclavian steal syndrome, periventricular leukomalacia, vascular headache, cluster headache, migraine, and vertebrobasilar insufficiency.

[0602] Embolisms include, but are not limited to, air embolisms, amniotic fluid embolisms, cholesterol embolisms, blue toe syndrome, fat embolisms, pulmonary embolisms, and thromboembolisms. Thrombosis include, but are not limited to, coronary thrombosis, hepatic vein thrombosis, retinal vein occlusion, carotid artery thrombosis, sinus thrombosis, Wallenberg's syndrome, and thrombophlebitis.

tamponade, endocarditis (including bacterial), heart aneurysm, cardiac arrest, congestive heart failure, congestive cardiomyopathy, paroxysmal dyspnea, cardiac edema, heart hypertrophy, congestive cardiomyopathy, left ventricular hypertrophy, right ventricular hypertrophy, post-infarction heart rupture, ventricular septal rupture, heart valve diseases, myocardial diseases, myocardial ischemia, pericardial effusion, pericarditis (including constrictive and tuberculous), pneumopericardium, postpericardiotomy syndrome, pulmonary heart disease, rheumatic heart disease, ventricular dysfunction, hyperemia, cardiovascular pregnancy complications, Scimitar Syndrome, cardiovascular syphilis, and cardiovascular tuberculosis.

[0594] Arrhythmias include, but are not limited to, sinus arrhythmia, atrial fibrillation, atrial flutter, bradycardia, extrasystole, Adams-Stokes Syndrome, bundle-branch block, sinoatrial block, long QT syndrome, parasystole, Lown-Ganong-Levine Syndrome, Mahaim-type pre-excitation syndrome, Wolff-Parkinson-White syndrome, sick sinus syndrome, tachycardias, and ventricular fibrillation. Tachycardias include paroxysmal tachycardia, supraventricular tachycardia, accelerated idioventricular rhythm, atrioventricular nodal reentry tachycardia, ectopic atrial tachycardia, ectopic junctional tachycardia, sinoatrial nodal reentry tachycardia, sinus tachycardia, Torsades de Pointes, and ventricular tachycardia.

[0595] Heart valve diseases include, but are not limited to, aortic valve insufficiency, aortic valve stenosis, heart murmurs, aortic valve prolapse, mitral valve prolapse, tricuspid valve prolapse, mitral valve insufficiency, mitral valve stenosis, pulmonary atresia, pulmonary valve insufficiency, pulmonary valve stenosis, tricuspid atresia, tricuspid valve insufficiency, and tricuspid valve stenosis.

[0596] Myocardial diseases include, but are not limited to, alcoholic cardiomyopathy, congestive cardiomyopathy, hypertrophic cardiomyopathy, aortic subvalvular stenosis, pulmonary subvalvular stenosis, restrictive cardiomyopathy, Chagas cardiomyopathy, endocardial fibroelastosis, endomyocardial fibrosis, Kearns Syndrome, myocardial reperfusion injury, and myocarditis.

[0597] Myocardial ischemias include, but are not limited to, coronary disease, such as angina pectoris, coronary aneurysm, coronary arteriosclerosis, coronary thrombosis, coronary vasospasm, myocardial infarction and myocardial stunning.

nephropathy, focal segmental glomerulosclerosis (FSGS), necrotizing glomerulonephritis, and renal papillary necrosis), cancers of the kidney (e.g., nephroma, hypernephroma, nephroblastoma, renal cell cancer, transitional cell cancer, renal adenocarcinoma, squamous cell cancer, and Wilm's tumor), and electrolyte imbalances (e.g., nephrocalcinosis, pyuria, edema, hydronephritis, proteinuria, hyponatremia, hypernatremia, hypokalemia, hyperkalemia, hypocalcemia, hypercalcemia, hypophosphatemia, and hyperphosphatemia).

[0590] Polypeptides may be administered using any method known in the art, including, but not limited to, direct needle injection at the delivery site, intravenous injection, topical administration, catheter infusion, biolistic injectors, particle accelerators, gelfoam sponge depots, other commercially available depot materials, osmotic pumps, oral or suppository solid pharmaceutical formulations, decanting or topical applications during surgery, aerosol delivery. Such methods are known in the art. Polypeptides may be administered as part of a Therapeutic, described in more detail below. Methods of delivering polynucleotides are described in more detail herein.

Cardiovascular Disorders

[0591] Polynucleotides or polypeptides, or agonists or antagonists of the present invention, may be used to treat, prevent, diagnose, and/or prognose cardiovascular disorders, including, but not limited to, peripheral artery disease, such as limb ischemia.

[0592] Cardiovascular disorders include, but are not limited to, cardiovascular abnormalities, such as arterio-arterial fistula, arteriovenous fistula, cerebral arteriovenous malformations, congenital heart defects, pulmonary atresia, and Scimitar Syndrome. Congenital heart defects include, but are not limited to, aortic coarctation, cor triatriatum, coronary vessel anomalies, crisscross heart, dextrocardia, patent ductus arteriosus, Ebstein's anomaly, Eisenmenger complex, hypoplastic left heart syndrome, levocardia, tetralogy of fallot, transposition of great vessels, double outlet right ventricle, tricuspid atresia, persistent truncus arteriosus, and heart septal defects, such as aortopulmonary septal defect, endocardial cushion defects, Lutembacher's Syndrome, trilogy of Fallot, ventricular heart septal defects.

[0593] Cardiovascular disorders also include, but are not limited to, heart disease, such as arrhythmias, carcinoid heart disease, high cardiac output, low cardiac output, cardiac

nephritis, blood vessel disorders of kidney, metabolic and congenital kidney disorders, urinary disorders of the kidney, autoimmune disorders, sclerosis and necrosis, electrolyte imbalance, and kidney cancers.

[0587] Kidney diseases which can be diagnosed, prognosed, prevented, and/or treated with compositions of the invention include, but are not limited to, acute kidney failure, chronic kidney failure, atheroembolic renal failure, end-stage renal disease, inflammatory diseases of the kidney (e.g., acute glomerulonephritis, postinfectious glomerulonephritis, rapidly progressive glomerulonephritis, nephrotic syndrome, membranous glomerulonephritis, familial nephrotic syndrome, membranoproliferative glomerulonephritis I and II, mesangial proliferative glomerulonephritis, chronic glomerulonephritis, acute tubulointerstitial nephritis, chronic tubulointerstitial nephritis, acute post-streptococcal glomerulonephritis (PSGN), pyelonephritis, lupus nephritis, chronic nephritis, interstitial nephritis, and post-streptococcal glomerulonephritis), blood vessel disorders of the kidneys (e.g., kidney infarction, atheroembolic kidney disease, cortical necrosis, malignant nephrosclerosis, renal vein thrombosis, renal underperfusion, renal retinopathy, renal ischemia-reperfusion, renal artery embolism, and renal artery stenosis), and kidney disorders resulting from urinary tract disease (e.g., pyelonephritis, hydronephrosis, urolithiasis (renal lithiasis, nephrolithiasis), reflux nephropathy, urinary tract infections, urinary retention, and acute or chronic unilateral obstructive uropathy.)

[0588] In addition, compositions of the invention can be used to diagnose, prognose, prevent, and/or treat metabolic and congenital disorders of the kidney (e.g., uremia, renal amyloidosis, renal osteodystrophy, renal tubular acidosis, renal glycosuria, nephrogenic diabetes insipidus, cystinuria, Fanconi's syndrome, renal fibrocystic osteosis (renal rickets), Hartnup disease, Bartter's syndrome, Liddle's syndrome, polycystic kidney disease, medullary cystic disease, medullary sponge kidney, Alport's syndrome, nail-patella syndrome, congenital nephrotic syndrome, CRUSH syndrome, horseshoe kidney, diabetic nephropathy, nephrogenic diabetes insipidus, analgesic nephropathy, kidney stones, and membranous nephropathy), and autoimmune disorders of the kidney (e.g., systemic lupus erythematosus (SLE), Goodpasture syndrome, IgA nephropathy, and IgM mesangial proliferative glomerulonephritis).

[0589] Compositions of the invention can also be used to diagnose, prognose, prevent, and/or treat sclerotic or necrotic disorders of the kidney (e.g., glomerulosclerosis, diabetic

small molecule drugs or adjuvants, such as apoptonin, galectins, thioredoxins, anti-inflammatory proteins (See for example, *Mutat Res* 400(1-2):447-55 (1998), *Med Hypotheses*.50(5):423-33 (1998), *Chem Biol Interact.* Apr 24;111-112:23-34 (1998), *J Mol Med*.76(6):402-12 (1998), *Int J Tissue React*;20(1):3-15 (1998), which are all hereby incorporated by reference).

[0583] Polypeptides, including protein fusions to, or fragments thereof, of the present invention are useful in inhibiting the metastasis of proliferative cells or tissues. Inhibition may occur as a direct result of administering polypeptides, or antibodies directed to said polypeptides as described elsewhere herein, or indirectly, such as activating the expression of proteins known to inhibit metastasis, for example alpha 4 integrins, (See, e.g., *Curr Top Microbiol Immunol* 1998;231:125-41, which is hereby incorporated by reference). Such therapeutic affects of the present invention may be achieved either alone, or in combination with small molecule drugs or adjuvants.

[0584] In another embodiment, the invention provides a method of delivering compositions containing the polypeptides of the invention (e.g., compositions containing polypeptides or polypeptide antibodies associated with heterologous polypeptides, heterologous nucleic acids, toxins, or prodrugs) to targeted cells expressing the polypeptide of the present invention. Polypeptides or polypeptide antibodies of the invention may be associated with with heterologous polypeptides, heterologous nucleic acids, toxins, or prodrugs via hydrophobic, hydrophilic, ionic and/or covalent interactions.

[0585] Polypeptides, protein fusions to, or fragments thereof, of the present invention are useful in enhancing the immunogenicity and/or antigenicity of proliferating cells or tissues, either directly, such as would occur if the polypeptides of the present invention 'vaccinated' the immune response to respond to proliferative antigens and immunogens, or indirectly, such as in activating the expression of proteins known to enhance the immune response (e.g. chemokines), to said antigens and immunogens.

Renal Disorders

[0586] Polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention, may be used to treat, prevent, diagnose, and/or prognose disorders of the renal system. Renal disorders which can be diagnosed, prognosed, prevented, and/or treated with compositions of the invention include, but are not limited to, kidney failure,

growth factors, for example., which serve to increase the number or activity of effector cells which interact with the antibodies.

[0580] It is preferred to use high affinity and/or potent *in vivo* inhibiting and/or neutralizing antibodies against polypeptides or polynucleotides of the present invention, fragments or regions thereof, for both immunoassays directed to and therapy of disorders related to polynucleotides or polypeptides, including fragments thereof, of the present invention. Such antibodies, fragments, or regions, will preferably have an affinity for polynucleotides or polypeptides, including fragments thereof. Preferred binding affinities include those with a dissociation constant or K_d less than $5 \times 10^{-6}M$, $10^{-6}M$, $5 \times 10^{-7}M$, $10^{-7}M$, $5 \times 10^{-8}M$, $10^{-8}M$, $5 \times 10^{-9}M$, $10^{-9}M$, $5 \times 10^{-10}M$, $10^{-10}M$, $5 \times 10^{-11}M$, $10^{-11}M$, $5 \times 10^{-12}M$, $10^{-12}M$, $5 \times 10^{-13}M$, $10^{-13}M$, $5 \times 10^{-14}M$, $10^{-14}M$, $5 \times 10^{-15}M$, and $10^{-15}M$.

[0581] Moreover, polypeptides of the present invention are useful in inhibiting the angiogenesis of proliferative cells or tissues, either alone, as a protein fusion, or in combination with other polypeptides directly or indirectly, as described elsewhere herein. In a most preferred embodiment, said anti-angiogenesis effect may be achieved indirectly, for example, through the inhibition of hematopoietic, tumor-specific cells, such as tumor-associated macrophages (See Joseph IB, et al. J Natl Cancer Inst, 90(21):1648-53 (1998), which is hereby incorporated by reference). Antibodies directed to polypeptides or polynucleotides of the present invention may also result in inhibition of angiogenesis directly, or indirectly (See Witte L, et al., Cancer Metastasis Rev. 17(2):155-61 (1998), which is hereby incorporated by reference)).

[0582] Polypeptides, including protein fusions, of the present invention, or fragments thereof may be useful in inhibiting proliferative cells or tissues through the induction of apoptosis. Said polypeptides may act either directly, or indirectly to induce apoptosis of proliferative cells and tissues, for example in the activation of a death-domain receptor, such as tumor necrosis factor (TNF) receptor-1, CD95 (Fas/APO-1), TNF-receptor-related apoptosis-mediated protein (TRAMP) and TNF-related apoptosis-inducing ligand (TRAIL) receptor-1 and -2 (See Schulze-Osthoff K, et.al., Eur J Biochem 254(3):439-59 (1998), which is hereby incorporated by reference). Moreover, in another preferred embodiment of the present invention, said polypeptides may induce apoptosis through other mechanisms, such as in the activation of other proteins which will activate apoptosis, or through stimulating the expression of said proteins, either alone or in combination with

characterized by single or multiple local abnormal proliferations of cells, groups of cells, or tissues, whether benign or malignant.

[0575] Any amount of the polynucleotides of the present invention may be administered as long as it has a biologically inhibiting effect on the proliferation of the treated cells. Moreover, it is possible to administer more than one of the polynucleotide of the present invention simultaneously to the same site. By "biologically inhibiting" is meant partial or total growth inhibition as well as decreases in the rate of proliferation or growth of the cells. The biologically inhibitory dose may be determined by assessing the effects of the polynucleotides of the present invention on target malignant or abnormally proliferating cell growth in tissue culture, tumor growth in animals and cell cultures, or any other method known to one of ordinary skill in the art.

[0576] The present invention is further directed to antibody-based therapies which involve administering of anti-polypeptides and anti-polynucleotide antibodies to a mammalian, preferably human, patient for treating one or more of the described disorders. Methods for producing anti-polypeptides and anti-polynucleotide antibodies polyclonal and monoclonal antibodies are described in detail elsewhere herein. Such antibodies may be provided in pharmaceutically acceptable compositions as known in the art or as described herein.

[0577] A summary of the ways in which the antibodies of the present invention may be used therapeutically includes binding polynucleotides or polypeptides of the present invention locally or systemically in the body or by direct cytotoxicity of the antibody, e.g. as mediated by complement (CDC) or by effector cells (ADCC). Some of these approaches are described in more detail below. Armed with the teachings provided herein, one of ordinary skill in the art will know how to use the antibodies of the present invention for diagnostic, monitoring or therapeutic purposes without undue experimentation.

[0578] In particular, the antibodies, fragments and derivatives of the present invention are useful for treating a subject having or developing cell proliferative and/or differentiation disorders as described herein. Such treatment comprises administering a single or multiple doses of the antibody, or a fragment, derivative, or a conjugate thereof.

[0579] The antibodies of this invention may be advantageously utilized in combination with other monoclonal or chimeric antibodies, or with lymphokines or hematopoietic

[0571] Polynucleotides of the present invention may be useful in repressing expression of oncogenic genes or antigens. By "repressing expression of the oncogenic genes " is intended the suppression of the transcription of the gene, the degradation of the gene transcript (pre-message RNA), the inhibition of splicing, the destruction of the messenger RNA, the prevention of the post-translational modifications of the protein, the destruction of the protein, or the inhibition of the normal function of the protein.

[0572] For local administration to abnormally proliferating cells, polynucleotides of the present invention may be administered by any method known to those of skill in the art including, but not limited to transfection, electroporation, microinjection of cells, or in vehicles such as liposomes, lipofectin, or as naked polynucleotides, or any other method described throughout the specification. The polynucleotide of the present invention may be delivered by known gene delivery systems such as, but not limited to, retroviral vectors (Gilboa, J. Virology 44:845 (1982); Hocke, Nature 320:275 (1986); Wilson, et al., Proc. Natl. Acad. Sci. U.S.A. 85:3014), vaccinia virus system (Chakrabarty et al., Mol. Cell Biol. 5:3403 (1985) or other efficient DNA delivery systems (Yates et al., Nature 313:812 (1985)) known to those skilled in the art. These references are exemplary only and are hereby incorporated by reference. In order to specifically deliver or transfect cells which are abnormally proliferating and spare non-dividing cells, it is preferable to utilize a retrovirus, or adenoviral (as described in the art and elsewhere herein) delivery system known to those of skill in the art. Since host DNA replication is required for retroviral DNA to integrate and the retrovirus will be unable to self replicate due to the lack of the retrovirus genes needed for its life cycle. Utilizing such a retroviral delivery system for polynucleotides of the present invention will target said gene and constructs to abnormally proliferating cells and will spare the non-dividing normal cells.

[0573] The polynucleotides of the present invention may be delivered directly to cell proliferative disorder/disease sites in internal organs, body cavities and the like by use of imaging devices used to guide an injecting needle directly to the disease site. The polynucleotides of the present invention may also be administered to disease sites at the time of surgical intervention.

[0574] By "cell proliferative disease" is meant any human or animal disease or disorder, affecting any one or any combination of organs, cavities, or body parts, which is

[0567] Similarly, other hyperproliferative disorders can also be diagnosed, prognosed, prevented, and/or treated by polynucleotides, polypeptides, and/or agonists or antagonists of the invention. Examples of such hyperproliferative disorders include, but are not limited to: hypergammaglobulinemia, lymphoproliferative disorders, paraproteinemias, purpura, sarcoidosis, Sezary Syndrome, Waldenstrom's macroglobulinemia, Gaucher's Disease, histiocytosis, and any other hyperproliferative disease, besides neoplasia, located in an organ system listed above.

[0568] Another preferred embodiment utilizes polynucleotides of the present invention to inhibit aberrant cellular division, by gene therapy using the present invention, and/or protein fusions or fragments thereof.

[0569] Thus, the present invention provides a method for treating cell proliferative disorders by inserting into an abnormally proliferating cell a polynucleotide of the present invention, wherein said polynucleotide represses said expression.

[0570] Another embodiment of the present invention provides a method of treating cell-proliferative disorders in individuals comprising administration of one or more active gene copies of the present invention to an abnormally proliferating cell or cells. In a preferred embodiment, polynucleotides of the present invention is a DNA construct comprising a recombinant expression vector effective in expressing a DNA sequence encoding said polynucleotides. In another preferred embodiment of the present invention, the DNA construct encoding the polynucleotides of the present invention is inserted into cells to be treated utilizing a retrovirus, or more preferably an adenoviral vector (See G J. Nabel, et. al., PNAS 1999 96: 324-326, which is hereby incorporated by reference). In a most preferred embodiment, the viral vector is defective and will not transform non-proliferating cells, only proliferating cells. Moreover, in a preferred embodiment, the polynucleotides of the present invention inserted into proliferating cells either alone, or in combination with or fused to other polynucleotides, can then be modulated via an external stimulus (i.e. magnetic, specific small molecule, chemical, or drug administration, etc.), which acts upon the promoter upstream of said polynucleotides to induce expression of the encoded protein product. As such the beneficial therapeutic affect of the present invention may be expressly modulated (i.e. to increase, decrease, or inhibit expression of the present invention) based upon said external stimulus.

lymphangiosarcoma, lymphangioendotheliosarcoma, synovioma, mesothelioma, Ewing's tumor, leiomyosarcoma, rhabdomyosarcoma, colon carcinoma, pancreatic cancer, breast cancer, ovarian cancer, prostate cancer, squamous cell carcinoma, basal cell carcinoma, adenocarcinoma, sweat gland carcinoma, sebaceous gland carcinoma, papillary carcinoma, papillary adenocarcinomas, cystadenocarcinoma, medullary carcinoma, bronchogenic carcinoma, renal cell carcinoma, hepatoma, bile duct carcinoma, choriocarcinoma, seminoma, embryonal carcinoma, Wilm's tumor, cervical cancer, testicular tumor, lung carcinoma, small cell lung carcinoma, bladder carcinoma, epithelial carcinoma, glioma, astrocytoma, medulloblastoma, craniopharyngioma, ependymoma, pinealoma, emangioblastoma, acoustic neuroma, oligodendroglioma, meningioma, melanoma, neuroblastoma, and retinoblastoma.

[0565] Diseases associated with increased apoptosis that could be diagnosed, prognosed, prevented, and/or treated by polynucleotides, polypeptides, and/or agonists or antagonists of the invention, include AIDS; neurodegenerative disorders (such as Alzheimer's disease, Parkinson's disease, amyotrophic lateral sclerosis, retinitis pigmentosa, cerebellar degeneration and brain tumor or prior associated disease); autoimmune disorders (such as, multiple sclerosis, Sjogren's syndrome, Hashimoto's thyroiditis, biliary cirrhosis, Behcet's disease, Crohn's disease, polymyositis, systemic lupus erythematosus and immune-related glomerulonephritis and rheumatoid arthritis) myelodysplastic syndromes (such as aplastic anemia), graft v. host disease, ischemic injury (such as that caused by myocardial infarction, stroke and reperfusion injury), liver injury (e.g., hepatitis related liver injury, ischemia/reperfusion injury, cholestasis (bile duct injury) and liver cancer); toxin-induced liver disease (such as that caused by alcohol), septic shock, cachexia and anorexia.

[0566] Hyperproliferative diseases and/or disorders that could be diagnosed, prognosed, prevented, and/or treated by polynucleotides, polypeptides, and/or agonists or antagonists of the invention, include, but are not limited to, neoplasms located in the liver, abdomen, bone, breast, digestive system, pancreas, peritoneum, endocrine glands (adrenal, parathyroid, pituitary, testicles, ovary, thymus, thyroid), eye, head and neck, nervous system (central and peripheral), lymphatic system, pelvis, skin, soft tissue, spleen, thorax, and urogenital tract.

[0562] Additionally, polynucleotides, polypeptides, and/or agonists or antagonists of the invention may affect apoptosis, and therefore, would be useful in treating a number of diseases associated with increased cell survival or the inhibition of apoptosis. For example, diseases associated with increased cell survival or the inhibition of apoptosis that could be diagnosed, prognosed, prevented, and/or treated by polynucleotides, polypeptides, and/or agonists or antagonists of the invention, include cancers (such as follicular lymphomas, carcinomas with p53 mutations, and hormone-dependent tumors, including, but not limited to colon cancer, cardiac tumors, pancreatic cancer, melanoma, retinoblastoma, glioblastoma, lung cancer, intestinal cancer, testicular cancer, stomach cancer, neuroblastoma, myxoma, myoma, lymphoma, endothelioma, osteoblastoma, osteoclastoma, osteosarcoma, chondrosarcoma, adenoma, breast cancer, prostate cancer, Kaposi's sarcoma and ovarian cancer); autoimmune disorders such as, multiple sclerosis, Sjogren's syndrome, Hashimoto's thyroiditis, biliary cirrhosis, Behcet's disease, Crohn's disease, polymyositis, systemic lupus erythematosus and immune-related glomerulonephritis and rheumatoid arthritis) and viral infections (such as herpes viruses, pox viruses and adenoviruses), inflammation, graft v. host disease, acute graft rejection, and chronic graft rejection.

[0563] In preferred embodiments, polynucleotides, polypeptides, and/or agonists or antagonists of the invention are used to inhibit growth, progression, and/or metastasis of cancers, in particular those listed above.

[0564] Additional diseases or conditions associated with increased cell survival that could be diagnosed, prognosed, prevented, and/or treated by polynucleotides, polypeptides, and/or agonists or antagonists of the invention, include, but are not limited to, progression, and/or metastases of malignancies and related disorders such as leukemia (including acute leukemias (e.g., acute lymphocytic leukemia, acute myelocytic leukemia (including myeloblastic, promyelocytic, myelomonocytic, monocytic, and erythroleukemia)) and chronic leukemias (e.g., chronic myelocytic (granulocytic) leukemia and chronic lymphocytic leukemia)), polycythemia vera, lymphomas (e.g., Hodgkin's disease and non-Hodgkin's disease), multiple myeloma, Waldenstrom's macroglobulinemia, heavy chain disease, and solid tumors including, but not limited to, sarcomas and carcinomas such as fibrosarcoma, myxosarcoma, liposarcoma, chondrosarcoma, osteogenic sarcoma, chordoma, angiosarcoma, endotheliosarcoma,

dysplasia, congenital ectodermal dysplasia, craniodiaphysial dysplasia, craniocarpotarsal dysplasia, craniometaphysial dysplasia, dentin dysplasia, diaphysial dysplasia, ectodermal dysplasia, enamel dysplasia, encephalo-ophthalmic dysplasia, dysplasia epiphysialis hemimelia, dysplasia epiphysialis multiplex, dysplasia epiphysialis punctata, epithelial dysplasia, faciodigitogenital dysplasia, familial fibrous dysplasia of jaws, familial white folded dysplasia, fibromuscular dysplasia, fibrous dysplasia of bone, florid osseous dysplasia, hereditary renal-retinal dysplasia, hidrotic ectodermal dysplasia, hypohidrotic ectodermal dysplasia, lymphopenic thymic dysplasia, mammary dysplasia, mandibulofacial dysplasia, metaphysial dysplasia, Mondini dysplasia, monostotic fibrous dysplasia, mucoepithelial dysplasia, multiple epiphysial dysplasia, oculoauriculovertebral dysplasia, oculodentodigital dysplasia, oculovertbral dysplasia, odontogenic dysplasia, ophthalmomandibulomelic dysplasia, periapical cemental dysplasia, polyostotic fibrous dysplasia, pseudoachondroplastic spondyloepiphysial dysplasia, retinal dysplasia, septo-optic dysplasia, spondyloepiphysial dysplasia, and ventriculoradial dysplasia.

[0559] Additional pre-neoplastic disorders which can be diagnosed, prognosed, prevented, and/or treated with compositions of the invention (including polynucleotides, polypeptides, agonists or antagonists) include, but are not limited to, benign dysproliferative disorders (e.g., benign tumors, fibrocystic conditions, tissue hypertrophy, intestinal polyps, colon polyps, and esophageal dysplasia), leukoplakia, keratoses, Bowen's disease, Farmer's Skin, solar cheilitis, and solar keratosis.

[0560] In another embodiment, a polypeptide of the invention, or polynucleotides, antibodies, agonists, or antagonists corresponding to that polypeptide, may be used to diagnose and/or prognose disorders associated with the tissue(s) in which the polypeptide of the invention is expressed, including one, two, three, four, five, or more tissues disclosed in Table 1, column 8 (Tissue Distribution Library Code).

[0561] In another embodiment, polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention conjugated to a toxin or a radioactive isotope, as described herein, may be used to treat cancers and neoplasms, including, but not limited to those described herein. In a further preferred embodiment, polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention conjugated to a toxin or a radioactive isotope, as described herein, may be used to treat acute myelogenous leukemia.

Hyperplastic disorders which can be diagnosed, prognosed, prevented, and/or treated with compositions of the invention (including polynucleotides, polypeptides, agonists or antagonists) include, but are not limited to, angiofollicular mediastinal lymph node hyperplasia, angiolymphoid hyperplasia with eosinophilia, atypical melanocytic hyperplasia, basal cell hyperplasia, benign giant lymph node hyperplasia, cementum hyperplasia, congenital adrenal hyperplasia, congenital sebaceous hyperplasia, cystic hyperplasia, cystic hyperplasia of the breast, denture hyperplasia, ductal hyperplasia, endometrial hyperplasia, fibromuscular hyperplasia, focal epithelial hyperplasia, gingival hyperplasia, inflammatory fibrous hyperplasia, inflammatory papillary hyperplasia, intravascular papillary endothelial hyperplasia, nodular hyperplasia of prostate, nodular regenerative hyperplasia, pseudoepitheliomatous hyperplasia, senile sebaceous hyperplasia, and verrucous hyperplasia.

[0557] Metaplasia is a form of controlled cell growth in which one type of adult or fully differentiated cell substitutes for another type of adult cell. Metaplastic disorders which can be diagnosed, prognosed, prevented, and/or treated with compositions of the invention (including polynucleotides, polypeptides, agonists or antagonists) include, but are not limited to, agnogenic myeloid metaplasia, apocrine metaplasia, atypical metaplasia, autoparenchymatous metaplasia, connective tissue metaplasia, epithelial metaplasia, intestinal metaplasia, metaplastic anemia, metaplastic ossification, metaplastic polyps, myeloid metaplasia, primary myeloid metaplasia, secondary myeloid metaplasia, squamous metaplasia, squamous metaplasia of amnion, and symptomatic myeloid metaplasia.

[0558] Dysplasia is frequently a forerunner of cancer, and is found mainly in the epithelia; it is the most disorderly form of non-neoplastic cell growth, involving a loss in individual cell uniformity and in the architectural orientation of cells. Dysplastic cells often have abnormally large, deeply stained nuclei, and exhibit pleomorphism. Dysplasia characteristically occurs where there exists chronic irritation or inflammation. Dysplastic disorders which can be diagnosed, prognosed, prevented, and/or treated with compositions of the invention (including polynucleotides, polypeptides, agonists or antagonists) include, but are not limited to, anhidrotic ectodermal dysplasia, anterofacial dysplasia, asphyxiating thoracic dysplasia, atriadigital dysplasia, bronchopulmonary dysplasia, cerebral dysplasia, cervical dysplasia, chondroectodermal dysplasia, cleidocranial

Myeloma/Plasma Cell Neoplasm, Myelodysplastic Syndrome, Myelogenous Leukemia, Myeloid Leukemia, Myeloproliferative Disorders, Nasal Cavity and Paranasal Sinus Cancer, Nasopharyngeal Cancer, Neuroblastoma, Non-Hodgkin's Lymphoma During Pregnancy, Nonmelanoma Skin Cancer, Non-Small Cell Lung Cancer, Occult Primary Metastatic Squamous Neck Cancer, Oropharyngeal Cancer, Osteo-/Malignant Fibrous Sarcoma, Osteosarcoma/Malignant Fibrous Histiocytoma, Osteosarcoma/Malignant Fibrous Histiocytoma of Bone, Ovarian Epithelial Cancer, Ovarian Germ Cell Tumor, Ovarian Low Malignant Potential Tumor, Pancreatic Cancer, Paraproteinemias, Purpura, Parathyroid Cancer, Penile Cancer, Pheochromocytoma, Pituitary Tumor, Plasma Cell Neoplasm/Multiple Myeloma, Primary Central Nervous System Lymphoma, Primary Liver Cancer, Prostate Cancer, Rectal Cancer, Renal Cell Cancer, Renal Pelvis and Ureter Cancer, Retinoblastoma, Rhabdomyosarcoma, Salivary Gland Cancer, Sarcoidosis Sarcomas, Sezary Syndrome, Skin Cancer, Small Cell Lung Cancer, Small Intestine Cancer, Soft Tissue Sarcoma, Squamous Neck Cancer, Stomach Cancer, Supratentorial Primitive Neuroectodermal and Pineal Tumors, T-Cell Lymphoma, Testicular Cancer, Thymoma, Thyroid Cancer, Transitional Cell Cancer of the Renal Pelvis and Ureter, Transitional Renal Pelvis and Ureter Cancer, Trophoblastic Tumors, Ureter and Renal Pelvis Cell Cancer, Urethral Cancer, Uterine Cancer, Uterine Sarcoma, Vaginal Cancer, Visual Pathway and Hypothalamic Glioma, Vulvar Cancer, Waldenstrom's Macroglobulinemia, Wilms' Tumor, and any other hyperproliferative disease, besides neoplasia, located in an organ system listed above.

[0555] In another preferred embodiment, polynucleotides or polypeptides, or agonists or antagonists of the present invention are used to diagnose, prognose, prevent, and/or treat premalignant conditions and to prevent progression to a neoplastic or malignant state, including but not limited to those disorders described above. Such uses are indicated in conditions known or suspected of preceding progression to neoplasia or cancer, in particular, where non-neoplastic cell growth consisting of hyperplasia, metaplasia, or most particularly, dysplasia has occurred (for review of such abnormal growth conditions, see Robbins and Angell, 1976, Basic Pathology, 2d Ed., W. B. Saunders Co., Philadelphia, pp. 68-79.)

[0556] Hyperplasia is a form of controlled cell proliferation, involving an increase in cell number in a tissue or organ, without significant alteration in structure or function.

(Primary) Hepatocellular Cancer, Adult (Primary) Liver Cancer, Adult Acute Lymphocytic Leukemia, Adult Acute Myeloid Leukemia, Adult Hodgkin's Disease, Adult Hodgkin's Lymphoma, Adult Lymphocytic Leukemia, Adult Non-Hodgkin's Lymphoma, Adult Primary Liver Cancer, Adult Soft Tissue Sarcoma, AIDS-Related Lymphoma, AIDS-Related Malignancies, Anal Cancer, Astrocytoma, Bile Duct Cancer, Bladder Cancer, Bone Cancer, Brain Stem Glioma, Brain Tumors, Breast Cancer, Cancer of the Renal Pelvis and Ureter, Central Nervous System (Primary) Lymphoma, Central Nervous System Lymphoma, Cerebellar Astrocytoma, Cerebral Astrocytoma, Cervical Cancer, Childhood (Primary) Hepatocellular Cancer, Childhood (Primary) Liver Cancer, Childhood Acute Lymphoblastic Leukemia, Childhood Acute Myeloid Leukemia, Childhood Brain Stem Glioma, Childhood Cerebellar Astrocytoma, Childhood Cerebral Astrocytoma, Childhood Extracranial Germ Cell Tumors, Childhood Hodgkin's Disease, Childhood Hodgkin's Lymphoma, Childhood Hypothalamic and Visual Pathway Glioma, Childhood Lymphoblastic Leukemia, Childhood Medulloblastoma, Childhood Non-Hodgkin's Lymphoma, Childhood Pineal and Supratentorial Primitive Neuroectodermal Tumors, Childhood Primary Liver Cancer, Childhood Rhabdomyosarcoma, Childhood Soft Tissue Sarcoma, Childhood Visual Pathway and Hypothalamic Glioma, Chronic Lymphocytic Leukemia, Chronic Myelogenous Leukemia, Colon Cancer, Cutaneous T-Cell Lymphoma, Endocrine Pancreas Islet Cell Carcinoma, Endometrial Cancer, Ependymoma, Epithelial Cancer, Esophageal Cancer, Ewing's Sarcoma and Related Tumors, Exocrine Pancreatic Cancer, Extracranial Germ Cell Tumor, Extragonadal Germ Cell Tumor, Extrahepatic Bile Duct Cancer, Eye Cancer, Female Breast Cancer, Gaucher's Disease, Gallbladder Cancer, Gastric Cancer, Gastrointestinal Carcinoid Tumor, Gastrointestinal Tumors, Germ Cell Tumors, Gestational Trophoblastic Tumor, Hairy Cell Leukemia, Head and Neck Cancer, Hepatocellular Cancer, Hodgkin's Disease, Hodgkin's Lymphoma, Hypergammaglobulinemia, Hypopharyngeal Cancer, Intestinal Cancers, Intraocular Melanoma, Islet Cell Carcinoma, Islet Cell Pancreatic Cancer, Kaposi's Sarcoma, Kidney Cancer, Laryngeal Cancer, Lip and Oral Cavity Cancer, Liver Cancer, Lung Cancer, Lymphoproliferative Disorders, Macroglobulinemia, Male Breast Cancer, Malignant Mesothelioma, Malignant Thymoma, Medulloblastoma, Melanoma, Mesothelioma, Metastatic Occult Primary Squamous Neck Cancer, Metastatic Primary Squamous Neck Cancer, Metastatic Squamous Neck Cancer, Multiple Myeloma, Multiple

[0549] In other embodiments, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful as an agent to increase cytokine production.

[0550] In other embodiments, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in preventing, diagnosing, and/or treating primary hematopoietic disorders.

Hyperproliferative Disorders

[0551] In certain embodiments, polynucleotides or polypeptides, or agonists or antagonists of the present invention can be used to treat or detect hyperproliferative disorders, including neoplasms. Polynucleotides or polypeptides, or agonists or antagonists of the present invention may inhibit the proliferation of the disorder through direct or indirect interactions. Alternatively, Polynucleotides or polypeptides, or agonists or antagonists of the present invention may proliferate other cells which can inhibit the hyperproliferative disorder.

[0552] For example, by increasing an immune response, particularly increasing antigenic qualities of the hyperproliferative disorder or by proliferating, differentiating, or mobilizing T-cells, hyperproliferative disorders can be treated. This immune response may be increased by either enhancing an existing immune response, or by initiating a new immune response. Alternatively, decreasing an immune response may also be a method of treating hyperproliferative disorders, such as a chemotherapeutic agent.

[0553] Examples of hyperproliferative disorders that can be treated or detected by polynucleotides or polypeptides, or agonists or antagonists of the present invention include, but are not limited to neoplasms located in the: colon, abdomen, bone, breast, digestive system, liver, pancreas, peritoneum, endocrine glands (adrenal, parathyroid, pituitary, testicles, ovary, thymus, thyroid), eye, head and neck, nervous (central and peripheral), lymphatic system, pelvis, skin, soft tissue, spleen, thorax, and urogenital tract.

[0554] Similarly, other hyperproliferative disorders can also be treated or detected by polynucleotides or polypeptides, or agonists or antagonists of the present invention. Examples of such hyperproliferative disorders include, but are not limited to: Acute Childhood Lymphoblastic Leukemia, Acute Lymphoblastic Leukemia, Acute Lymphocytic Leukemia, Acute Myeloid Leukemia, Adrenocortical Carcinoma, Adult

prognosing, preventing, and/or treating leukemias and lymphomas including, but not limited to, acute lymphocytic (lymphoblastic) leukemia (ALL), acute myeloid (myelocytic, myelogenous, myeloblastic, or myelomonocytic) leukemia, chronic lymphocytic leukemia (e.g., B cell leukemias, T cell leukemias, Sezary syndrome, and Hairy cell leukemia), chronic myelocytic (myeloid, myelogenous, or granulocytic) leukemia, Hodgkin's lymphoma, non-hodgkin's lymphoma, Burkitt's lymphoma, and mycosis fungoides.

[0544] In other embodiments, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in diagnosing, prognosing, preventing, and/or treating diseases and disorders of plasma cells including, but not limited to, plasma cell dyscrasias, monoclonal gammaopathies, monoclonal gammopathies of undetermined significance, multiple myeloma, macroglobulinemia, Waldenstrom's macroglobulinemia, cryoglobulinemia, and Raynaud's phenomenon.

[0545] In other embodiments, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in treating, preventing, and/or diagnosing myeloproliferative disorders, including but not limited to, polycythemia vera, relative polycythemia, secondary polycythemia, myelofibrosis, acute myelofibrosis, agnogenic myeloid metaplasia, thrombocythemia, (including both primary and secondary thrombocythemia) and chronic myelocytic leukemia.

[0546] In other embodiments, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful as a treatment prior to surgery, to increase blood cell production.

[0547] In other embodiments, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful as an agent to enhance the migration, phagocytosis, superoxide production, antibody dependent cellular cytotoxicity of neutrophils, eosinophils and macrophages.

[0548] In other embodiments, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful as an agent to increase the number of stem cells in circulation prior to stem cells pheresis. In another specific embodiment, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful as an agent to increase the number of stem cells in circulation prior to platelet pheresis.

be diagnosed, prognosed, prevented, and/or treated by the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention include, but are not limited to, infantile genetic agranulocytosis, familial neutropenia, cyclic neutropenia, neutropenias resulting from or associated with dietary deficiencies (e.g., vitamin B 12 deficiency or folic acid deficiency), neutropenias resulting from or associated with drug treatments (e.g., antibiotic regimens such as penicillin treatment, sulfonamide treatment, anticoagulant treatment, anticonvulsant drugs, anti-thyroid drugs, and cancer chemotherapy), and neutropenias resulting from increased neutrophil destruction that may occur in association with some bacterial or viral infections, allergic disorders, autoimmune diseases, conditions in which an individual has an enlarged spleen (e.g., Felty syndrome, malaria and sarcoidosis), and some drug treatment regimens.

[0540] The polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in diagnosing, prognosing, preventing, and/or treating lymphocytopenias (decreased numbers of B and/or T lymphocytes), including, but not limited lymphocytopenias resulting from or associated with stress, drug treatments (e.g., drug treatment with corticosteroids, cancer chemotherapies, and/or radiation therapies), AIDS infection and/or other diseases such as, for example, cancer, rheumatoid arthritis, systemic lupus erythematosus, chronic infections, some viral infections and/or hereditary disorders (e.g., DiGeorge syndrome, Wiskott-Aldrich Syndrome, severe combined immunodeficiency, ataxia telangiectasia).

[0541] The polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in diagnosing, prognosing, preventing, and/or treating diseases and disorders associated with macrophage numbers and/or macrophage function including, but not limited to, Gaucher's disease, Niemann-Pick disease, Letterer-Siwe disease and Hand-Schuller-Christian disease.

[0542] In another embodiment, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in diagnosing, prognosing, preventing, and/or treating diseases and disorders associated with eosinophil numbers and/or eosinophil function including, but not limited to, idiopathic hypereosinophilic syndrome, eosinophilia-myalgia syndrome, and Hand-Schuller-Christian disease.

[0543] In yet another embodiment, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in diagnosing,

thromboplastin time (PTT), the activated partial thromboplastin time (aPTT), the activated clotting time (ACT), the recalcified activated clotting time, or the Lee-White Clotting time.

[0537] Several diseases and a variety of drugs can cause platelet dysfunction. Thus, in a specific embodiment, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in diagnosing, prognosing, preventing, and/or treating acquired platelet dysfunction such as platelet dysfunction accompanying kidney failure, leukemia, multiple myeloma, cirrhosis of the liver, and systemic lupus erythematosus as well as platelet dysfunction associated with drug treatments, including treatment with aspirin, ticlopidine, nonsteroidal anti-inflammatory drugs (used for arthritis, pain, and sprains), and penicillin in high doses.

[0538] In another embodiment, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in diagnosing, prognosing, preventing, and/or treating diseases and disorders characterized by or associated with increased or decreased numbers of white blood cells. Leukopenia occurs when the number of white blood cells decreases below normal. Leukopenias include, but are not limited to, neutropenia and lymphocytopenia. An increase in the number of white blood cells compared to normal is known as leukocytosis. The body generates increased numbers of white blood cells during infection. Thus, leukocytosis may simply be a normal physiological parameter that reflects infection. Alternatively, leukocytosis may be an indicator of injury or other disease such as cancer. Leukocytoses, include but are not limited to, eosinophilia, and accumulations of macrophages. In specific embodiments, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in diagnosing, prognosing, preventing, and/or treating leukopenia. In other specific embodiments, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in diagnosing, prognosing, preventing, and/or treating leukocytosis.

[0539] Leukopenia may be a generalized decreased in all types of white blood cells, or may be a specific depletion of particular types of white blood cells. Thus, in specific embodiments, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in diagnosing, prognosing, preventing, and/or treating decreases in neutrophil numbers, known as neutropenia. Neutropenias that may

antagonists of the present invention may be useful in treating, preventing, and/or diagnosing anemias associated with diseases including but not limited to, anemias associated with systemic lupus erythematosus, cancers, lymphomas, chronic renal disease, and enlarged spleens. The polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in treating, preventing, and/or diagnosing anemias arising from drug treatments such as anemias associated with methyl dopa, dapsone, and/or sulfadiazine. Additionally, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in treating, preventing, and/or diagnosing anemias associated with abnormal red blood cell architecture including but not limited to, hereditary spherocytosis, hereditary elliptocytosis, glucose-6-phosphate dehydrogenase deficiency, and sickle cell anemia.

[0534] The polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in treating, preventing, and/or diagnosing hemoglobin abnormalities, (e.g., those associated with sickle cell anemia, hemoglobin C disease, hemoglobin S-C disease, and hemoglobin E disease). Additionally, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in diagnosing, prognosing, preventing, and/or treating thalassemias, including, but not limited to major and minor forms of alpha-thalassemia and beta-thalassemia.

[0535] In another embodiment, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in diagnosing, prognosing, preventing, and/or treating bleeding disorders including, but not limited to, thrombocytopenia (e.g., idiopathic thrombocytopenic purpura, and thrombotic thrombocytopenic purpura), Von Willebrand's disease, hereditary platelet disorders (e.g., storage pool disease such as Chediak-Higashi and Hermansky-Pudlak syndromes, thromboxane A₂ dysfunction, thromboasthenia, and Bernard-Soulier syndrome), hemolytic-uremic syndrome, hemophelias such as hemophilia A or Factor VII deficiency and Christmas disease or Factor IX deficiency, Hereditary Hemorrhagic Telangiectasia, also known as Rendu-Osler-Weber syndrome, allergic purpura (Henoch Schonlein purpura) and disseminated intravascular coagulation.

[0536] The effect of the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention on the clotting time of blood may be monitored using any of the clotting tests known in the art including, but not limited to, whole blood partial

expressed, including one, two, three, four, five, or more tissues disclosed in Table 1, column 8 (Tissue Distribution Library Code).

[0531] The polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be used to modulate hematopoietic activity (the formation of blood cells). For example, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be used to increase the quantity of all or subsets of blood cells, such as, for example, erythrocytes, lymphocytes (B or T cells), myeloid cells (e.g., basophils, eosinophils, neutrophils, mast cells, macrophages) and platelets. The ability to decrease the quantity of blood cells or subsets of blood cells may be useful in the prevention, detection, diagnosis and/or treatment of anemias and leukopenias described below. Alternatively, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be used to decrease the quantity of all or subsets of blood cells, such as, for example, erythrocytes, lymphocytes (B or T cells), myeloid cells (e.g., basophils, eosinophils, neutrophils, mast cells, macrophages) and platelets.. The ability to decrease the quantity of blood cells or subsets of blood cells may be useful in the prevention, detection, diagnosis and/or treatment of leukocytoses, such as, for example eosinophilia.

[0532] The polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be used to prevent, treat, or diagnose blood dyscrasia.

[0533] Anemias are conditions in which the number of red blood cells or amount of hemoglobin (the protein that carries oxygen) in them is below normal. Anemia may be caused by excessive bleeding, decreased red blood cell production, or increased red blood cell destruction (hemolysis). The polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in treating, preventing, and/or diagnosing anemias. Anemias that may be treated prevented or diagnosed by the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention include iron deficiency anemia, hypochromic anemia, microcytic anemia, chlorosis, hereditary sideroblastic anemia, idiopathic acquired sideroblastic anemia, red cell aplasia, megaloblastic anemia (e.g., pernicious anemia, (vitamin B12 deficiency) and folic acid deficiency anemia), aplastic anemia, hemolytic anemias (e.g., autoimmune hemolytic anemia, microangiopathic hemolytic anemia, and paroxysmal nocturnal hemoglobinuria). The polynucleotides, polypeptides, antibodies, and/or agonists or

Blood-Related Disorders

[0528] The polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be used to modulate hemostatic (the stopping of bleeding) or thrombolytic (clot dissolving) activity. For example, by increasing hemostatic or thrombolytic activity, polynucleotides or polypeptides, and/or agonists or antagonists of the present invention could be used to treat or prevent blood coagulation diseases, disorders, and/or conditions (e.g., afibrinogenemia, factor deficiencies, hemophilia), blood platelet diseases, disorders, and/or conditions (e.g., thrombocytopenia), or wounds resulting from trauma, surgery, or other causes. Alternatively, polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention that can decrease hemostatic or thrombolytic activity could be used to inhibit or dissolve clotting. These molecules could be important in the treatment or prevention of heart attacks (infarction), strokes, or scarring.

[0529] In specific embodiments, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be used to prevent, diagnose, prognose, and/or treat thrombosis, arterial thrombosis, venous thrombosis, thromboembolism, pulmonary embolism, atherosclerosis, myocardial infarction, transient ischemic attack, unstable angina. In specific embodiments, the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be used for the prevention of occlusion of saphenous grafts, for reducing the risk of periprocedural thrombosis as might accompany angioplasty procedures, for reducing the risk of stroke in patients with atrial fibrillation including nonrheumatic atrial fibrillation, for reducing the risk of embolism associated with mechanical heart valves and or mitral valves disease. Other uses for the polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention, include, but are not limited to, the prevention of occlusions in extracorporeal devices (e.g., intravascular canulas, vascular access shunts in hemodialysis patients, hemodialysis machines, and cardiopulmonary bypass machines).

[0530] In another embodiment, a polypeptide of the invention, or polynucleotides, antibodies, agonists, or antagonists corresponding to that polypeptide, may be used to prevent, diagnose, prognose, and/or treat diseases and disorders of the blood and/or blood forming organs associated with the tissue(s) in which the polypeptide of the invention is

myeloma, Burkitt's lymphoma, EBV-transformed diseases, and/or diseases and disorders described in the section entitled "Hyperproliferative Disorders" elsewhere herein.

[0523] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a therapy for decreasing cellular proliferation of Large B-cell Lymphomas.

[0524] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a means of decreasing the involvement of B cells and Ig associated with Chronic Myelogenous Leukemia.

[0525] In specific embodiments, the compositions of the invention are used as an agent to boost immunoresponsiveness among B cell immunodeficient individuals, such as, for example, an individual who has undergone a partial or complete splenectomy.

[0526] Antagonists of the invention include, for example, binding and/or inhibitory antibodies, antisense nucleic acids, ribozymes or soluble forms of the polypeptides of the present invention (e.g., Fc fusion protein; see, e.g., Example 9). Agonists of the invention include, for example, binding or stimulatory antibodies, and soluble forms of the polypeptides (e.g., Fc fusion proteins; see, e.g., Example 9). polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention may be employed in a composition with a pharmaceutically acceptable carrier, e.g., as described herein.

[0527] In another embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are administered to an animal (including, but not limited to, those listed above, and also including transgenic animals) incapable of producing functional endogenous antibody molecules or having an otherwise compromised endogenous immune system, but which is capable of producing human immunoglobulin molecules by means of a reconstituted or partially reconstituted immune system from another animal (see, e.g., published PCT Application Nos. WO98/24893, WO/9634096, WO/9633735, and WO/9110741). Administration of polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention to such animals is useful for the generation of monoclonal antibodies against the polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention.

[0520] In a specific embodiment, polynucleotides or polypeptides, and/or agonists thereof are used to diagnose, prognose, treat, and/or prevent a disorder characterized by primary or acquired immunodeficiency, deficient serum immunoglobulin production, recurrent infections, and/or immune system dysfunction. Moreover, polynucleotides or polypeptides, and/or agonists thereof may be used to treat or prevent infections of the joints, bones, skin, and/or parotid glands, blood-borne infections (e.g., sepsis, meningitis, septic arthritis, and/or osteomyelitis), autoimmune diseases (e.g., those disclosed herein), inflammatory disorders, and malignancies, and/or any disease or disorder or condition associated with these infections, diseases, disorders and/or malignancies) including, but not limited to, CVID, other primary immune deficiencies, HIV disease, CLL, recurrent bronchitis, sinusitis, otitis media, conjunctivitis, pneumonia, hepatitis, meningitis, herpes zoster (e.g., severe herpes zoster), and/or pneumocystis carinii. Other diseases and disorders that may be prevented, diagnosed, prognosed, and/or treated with polynucleotides or polypeptides, and/or agonists of the present invention include, but are not limited to, HIV infection, HTLV-BLV infection, lymphopenia, phagocyte bactericidal dysfunction anemia, thrombocytopenia, and hemoglobinuria.

[0521] In another embodiment, polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention are used to treat, and/or diagnose an individual having common variable immunodeficiency disease ("CVID"; also known as "acquired agammaglobulinemia" and "acquired hypogammaglobulinemia") or a subset of this disease.

[0522] In a specific embodiment, polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be used to diagnose, prognose, prevent, and/or treat cancers or neoplasms including immune cell or immune tissue-related cancers or neoplasms. Examples of cancers or neoplasms that may be prevented, diagnosed, or treated by polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention include, but are not limited to, acute myelogenous leukemia, chronic myelogenous leukemia, Hodgkin's disease, non-Hodgkin's lymphoma, acute lymphocytic anemia (ALL) Chronic lymphocyte leukemia, plasmacytomas, multiple

hypergammaglobulinemia evident in such diseases as monoclonal gammopathy of undetermined significance (MGUS), Waldenstrom's disease, related idiopathic monoclonal gammopathies, and plasmacytomas.

[0513] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention may be employed for instance to inhibit polypeptide chemotaxis and activation of macrophages and their precursors, and of neutrophils, basophils, B lymphocytes and some T-cell subsets, e.g., activated and CD8 cytotoxic T cells and natural killer cells, in certain autoimmune and chronic inflammatory and infective diseases. Examples of autoimmune diseases are described herein and include multiple sclerosis, and insulin-dependent diabetes.

[0514] The polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention may also be employed to treat idiopathic hyper-eosinophilic syndrome by, for example, preventing eosinophil production and migration.

[0515] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used to enhance or inhibit complement mediated cell lysis.

[0516] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used to enhance or inhibit antibody dependent cellular cytotoxicity.

[0517] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention may also be employed for treating atherosclerosis, for example, by preventing monocyte infiltration in the artery wall.

[0518] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention may be employed to treat adult respiratory distress syndrome (ARDS).

[0519] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention may be useful for stimulating wound and tissue repair, stimulating angiogenesis, and/or stimulating the repair of vascular or lymphatic diseases or disorders. Additionally, agonists and antagonists of the invention may be used to stimulate the regeneration of mucosal surfaces.

for genetically inherited disorders resulting in immuno-incompetence/immunodeficiency such as observed among SCID patients.

[0506] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a means of activating monocytes/macrophages to defend against parasitic diseases that effect monocytes such as Leishmania.

[0507] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a means of regulating secreted cytokines that are elicited by polypeptides of the invention.

[0508] In another embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used in one or more of the applications deccribed herein, as they may apply to veterinary medicine.

[0509] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a means of blocking various aspects of immune responses to foreign agents or self. Examples of diseases or conditions in which blocking of certain aspects of immune responses may be desired include autoimmune disorders such as lupus, and arthritis, as well as immunoresponsiveness to skin allergies, inflammation, bowel disease, injury and diseases/disorders associated with pathogens.

[0510] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a therapy for preventing the B cell proliferation and Ig secretion associated with autoimmune diseases such as idiopathic thrombocytopenic purpura, systemic lupus erythematosus and multiple sclerosis.

[0511] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a inhibitor of B and/or T cell migration in endothelial cells. This activity disrupts tissue architecture or cognate responses and is useful, for example in disrupting immune responses, and blocking sepsis.

[0512] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a therapy for chronic

[0500] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a regulator of antigen presentation by monocytes, dendritic cells, and/or B-cells. In one embodiment, polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention enhance antigen presentation or antagonizes antigen presentation in vitro or in vivo. Moreover, in related embodiments, said enhancement or antagonism of antigen presentation may be useful as an anti-tumor treatment or to modulate the immune system.

[0501] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an agent to direct an individual's immune system towards development of a humoral response (i.e. TH2) as opposed to a TH1 cellular response.

[0502] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a means to induce tumor proliferation and thus make it more susceptible to anti-neoplastic agents. For example, multiple myeloma is a slowly dividing disease and is thus refractory to virtually all anti-neoplastic regimens. If these cells were forced to proliferate more rapidly their susceptibility profile would likely change.

[0503] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a stimulator of B cell production in pathologies such as AIDS, chronic lymphocyte disorder and/or Common Variable Immunodeficiency.

[0504] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a therapy for generation and/or regeneration of lymphoid tissues following surgery, trauma or genetic defect. In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used in the pretreatment of bone marrow samples prior to transplant.

[0505] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a gene-based therapy

[0496] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an agent to boost immunoresponsiveness among aged populations and/or neonates.

[0497] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an immune system enhancer prior to, during, or after bone marrow transplant and/or other transplants (e.g., allogeneic or xenogeneic organ transplantation). With respect to transplantation, compositions of the invention may be administered prior to, concomitant with, and/or after transplantation. In a specific embodiment, compositions of the invention are administered after transplantation, prior to the beginning of recovery of T-cell populations. In another specific embodiment, compositions of the invention are first administered after transplantation after the beginning of recovery of T cell populations, but prior to full recovery of B cell populations.

[0498] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an agent to boost immunoresponsiveness among individuals having an acquired loss of B cell function. Conditions resulting in an acquired loss of B cell function that may be ameliorated or treated by administering the polypeptides, antibodies, polynucleotides and/or agonists or antagonists thereof, include, but are not limited to, HIV Infection, AIDS, bone marrow transplant, and B cell chronic lymphocytic leukemia (CLL).

[0499] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an agent to boost immunoresponsiveness among individuals having a temporary immune deficiency. Conditions resulting in a temporary immune deficiency that may be ameliorated or treated by administering the polypeptides, antibodies, polynucleotides and/or agonists or antagonists thereof, include, but are not limited to, recovery from viral infections (e.g., influenza), conditions associated with malnutrition, recovery from infectious mononucleosis, or conditions associated with stress, recovery from measles, recovery from blood transfusion, and recovery from surgery.

[0488] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an antigen for the generation of antibodies to inhibit or enhance immune mediated responses against polypeptides of the invention.

[0489] In one embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are administered to an animal (e.g., mouse, rat, rabbit, hamster, guinea pig, pigs, micro-pig, chicken, camel, goat, horse, cow, sheep, dog, cat, non-human primate, and human, most preferably human) to boost the immune system to produce increased quantities of one or more antibodies (e.g., IgG, IgA, IgM, and IgE), to induce higher affinity antibody production and immunoglobulin class switching (e.g., IgG, IgA, IgM, and IgE), and/or to increase an immune response.

[0490] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a stimulator of B cell responsiveness to pathogens.

[0491] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an activator of T cells.

[0492] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an agent that elevates the immune status of an individual prior to their receipt of immunosuppressive therapies.

[0493] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an agent to induce higher affinity antibodies.

[0494] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an agent to increase serum immunoglobulin concentrations.

[0495] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an agent to accelerate recovery of immunocompromised individuals.

influenza A and B, parainfluenza, measles, cytomegalovirus, rabies, Junin, Chikungunya, Rift Valley Fever, herpes simplex, and yellow fever.

[0484] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an adjuvant to enhance anti-bacterial or anti-fungal immune responses. Anti-bacterial or anti-fungal immune responses that may be enhanced using the compositions of the invention as an adjuvant, include bacteria or fungus and bacteria or fungus associated diseases or symptoms described herein or otherwise known in the art. In specific embodiments, the compositions of the invention are used as an adjuvant to enhance an immune response to a bacteria or fungus, disease, or symptom selected from the group consisting of: tetanus, Diphtheria, botulism, and meningitis type B.

[0485] In another specific embodiment, the compositions of the invention are used as an adjuvant to enhance an immune response to a bacteria or fungus, disease, or symptom selected from the group consisting of: *Vibrio cholerae*, *Mycobacterium leprae*, *Salmonella typhi*, *Salmonella paratyphi*, *Meissneria meningitidis*, *Streptococcus pneumoniae*, Group B streptococcus, *Shigella spp.*, Enterotoxigenic *Escherichia coli*, Enterohemorrhagic *E. coli*, and *Borrelia burgdorferi*.

[0486] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an adjuvant to enhance anti-parasitic immune responses. Anti-parasitic immune responses that may be enhanced using the compositions of the invention as an adjuvant, include parasite and parasite associated diseases or symptoms described herein or otherwise known in the art. In specific embodiments, the compositions of the invention are used as an adjuvant to enhance an immune response to a parasite. In another specific embodiment, the compositions of the invention are used as an adjuvant to enhance an immune response to *Plasmodium* (malaria) or *Leishmania*.

[0487] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention may also be employed to treat infectious diseases including silicosis, sarcoidosis, and idiopathic pulmonary fibrosis; for example, by preventing the recruitment and activation of mononuclear phagocytes.

[0480] In other embodiments, polypeptides, antibodies, or polynucleotides of the invention, and/or agonists or antagonists thereof, are useful to diagnose, prognose, prevent, and/or treat immune complex diseases, including, but not limited to, serum sickness, post streptococcal glomerulonephritis, polyarteritis nodosa, and immune complex-induced vasculitis.

[0481] Polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the invention can be used to treat, detect, and/or prevent infectious agents. For example, by increasing the immune response, particularly increasing the proliferation activation and/or differentiation of B and/or T cells, infectious diseases may be treated, detected, and/or prevented. The immune response may be increased by either enhancing an existing immune response, or by initiating a new immune response. Alternatively, polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may also directly inhibit the infectious agent (refer to section of application listing infectious agents, etc), without necessarily eliciting an immune response.

[0482] In another embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a vaccine adjuvant that enhances immune responsiveness to an antigen. In a specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an adjuvant to enhance tumor-specific immune responses.

[0483] In another specific embodiment, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as an adjuvant to enhance anti-viral immune responses. Anti-viral immune responses that may be enhanced using the compositions of the invention as an adjuvant, include virus and virus associated diseases or symptoms described herein or otherwise known in the art. In specific embodiments, the compositions of the invention are used as an adjuvant to enhance an immune response to a virus, disease, or symptom selected from the group consisting of: AIDS, meningitis, Dengue, EBV, and hepatitis (e.g., hepatitis B). In another specific embodiment, the compositions of the invention are used as an adjuvant to enhance an immune response to a virus, disease, or symptom selected from the group consisting of: HIV/AIDS, respiratory syncytial virus, Dengue, rotavirus, Japanese B encephalitis,

Grave's disease, systemic lupus erythematosus, diabetes mellitus, and allogenic transplant rejection).

[0478] Because inflammation is a fundamental defense mechanism, inflammatory disorders can effect virtually any tissue of the body. Accordingly, polynucleotides, polypeptides, and antibodies of the invention, as well as agonists or antagonists thereof, have uses in the treatment of tissue-specific inflammatory disorders, including, but not limited to, adrenalitis, alveolitis, angiocholecystitis, appendicitis, balanitis, blepharitis, bronchitis, bursitis, carditis, cellulitis, cervicitis, cholecystitis, chorditis, cochitis, colitis, conjunctivitis, cystitis, dermatitis, diverticulitis, encephalitis, endocarditis, esophagitis, eustachitis, fibrositis, folliculitis, gastritis, gastroenteritis, gingivitis, glossitis, hepatosplenitis, keratitis, labyrinthitis, laryngitis, lymphangitis, mastitis, media otitis, meningitis, metritis, mucitis, myocarditis, myositis, myringitis, nephritis, neuritis, orchitis, osteochondritis, otitis, pericarditis, peritendonitis, peritonitis, pharyngitis, phlebitis, poliomyelitis, prostatitis, pulpitis, retinitis, rhinitis, salpingitis, scleritis, sclerochoroiditis, scrotitis, sinusitis, spondylitis, steatitis, stomatitis, synovitis, syringitis, tendonitis, tonsillitis, urethritis, and vaginitis.

[0479] In specific embodiments, polypeptides, antibodies, or polynucleotides of the invention, and/or agonists or antagonists thereof, are useful to diagnose, prognose, prevent, and/or treat organ transplant rejections and graft-versus-host disease. Organ rejection occurs by host immune cell destruction of the transplanted tissue through an immune response. Similarly, an immune response is also involved in GVHD, but, in this case, the foreign transplanted immune cells destroy the host tissues. Polypeptides, antibodies, or polynucleotides of the invention, and/or agonists or antagonists thereof, that inhibit an immune response, particularly the activation, proliferation, differentiation, or chemotaxis of T-cells, may be an effective therapy in preventing organ rejection or GVHD. In specific embodiments, polypeptides, antibodies, or polynucleotides of the invention, and/or agonists or antagonists thereof, that inhibit an immune response, particularly the activation, proliferation, differentiation, or chemotaxis of T-cells, may be an effective therapy in preventing experimental allergic and hyperacute xenograft rejection.

using polypeptides, antibodies, or polynucleotides of the invention, and/or agonists or antagonists thereof. Moreover, these molecules can be used to treat, prevent, prognose, and/or diagnose anaphylaxis, hypersensitivity to an antigenic molecule, or blood group incompatibility.

[0476] Additionally, polypeptides or polynucleotides of the invention, and/or agonists or antagonists thereof, may be used to treat, prevent, diagnose and/or prognose IgE-mediated allergic reactions. Such allergic reactions include, but are not limited to, asthma, rhinitis, and eczema. In specific embodiments, polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be used to modulate IgE concentrations in vitro or in vivo.

[0477] Moreover, polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention have uses in the diagnosis, prognosis, prevention, and/or treatment of inflammatory conditions. For example, since polypeptides, antibodies, or polynucleotides of the invention, and/or agonists or antagonists of the invention may inhibit the activation, proliferation and/or differentiation of cells involved in an inflammatory response, these molecules can be used to prevent and/or treat chronic and acute inflammatory conditions. Such inflammatory conditions include, but are not limited to, for example, inflammation associated with infection (e.g., septic shock, sepsis, or systemic inflammatory response syndrome), ischemia-reperfusion injury, endotoxin lethality, complement-mediated hyperacute rejection, nephritis, cytokine or chemokine induced lung injury, inflammatory bowel disease, Crohn's disease, over production of cytokines (e.g., TNF or IL-1.), respiratory disorders (e.g., asthma and allergy); gastrointestinal disorders (e.g., inflammatory bowel disease); cancers (e.g., gastric, ovarian, lung, bladder, liver, and breast); CNS disorders (e.g., multiple sclerosis; ischemic brain injury and/or stroke, traumatic brain injury, neurodegenerative disorders (e.g., Parkinson's disease and Alzheimer's disease); AIDS-related dementia; and prion disease); cardiovascular disorders (e.g., atherosclerosis, myocarditis, cardiovascular disease, and cardiopulmonary bypass complications); as well as many additional diseases, conditions, and disorders that are characterized by inflammation (e.g., hepatitis, rheumatoid arthritis, gout, trauma, pancreatitis, sarcoidosis, dermatitis, renal ischemia-reperfusion injury,

[0470] In another specific preferred embodiment, systemic lupus erythematosus is treated, prevented, and/or diagnosed using polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention. In another specific preferred embodiment, idiopathic thrombocytopenia purpura is treated, prevented, and/or diagnosed using polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention.

[0471] In another specific preferred embodiment IgA nephropathy is treated, prevented, and/or diagnosed using polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention.

[0472] In a preferred embodiment, the autoimmune diseases and disorders and/or conditions associated with the diseases and disorders recited above are treated, prevented, diagnosed and/or prognosed using polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention.

[0473] In preferred embodiments, polypeptides, antibodies, polynucleotides and/or agonists or antagonists of the present invention are used as a immunosuppressive agent(s).

[0474] Polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in treating, preventing, prognosing, and/or diagnosing diseases, disorders, and/or conditions of hematopoietic cells. Polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention could be used to increase differentiation and proliferation of hematopoietic cells, including the pluripotent stem cells, in an effort to treat or prevent those diseases, disorders, and/or conditions associated with a decrease in certain (or many) types hematopoietic cells, including but not limited to, leukopenia, neutropenia, anemia, and thrombocytopenia. Alternatively, Polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention could be used to increase differentiation and proliferation of hematopoietic cells, including the pluripotent stem cells, in an effort to treat or prevent those diseases, disorders, and/or conditions associated with an increase in certain (or many) types of hematopoietic cells, including but not limited to, histiocytosis.

[0475] Allergic reactions and conditions, such as asthma (particularly allergic asthma) or other respiratory problems, may also be treated, prevented, diagnosed and/or prognosed

cell-mediated adrenal cytotoxicity, infertility (often characterized, e.g., by antispermatzoal antibodies), glomerulonephritis (often characterized, e.g., by glomerular basement membrane antibodies or immune complexes), bullous pemphigoid (often characterized, e.g., by IgG and complement in basement membrane), Sjogren's syndrome (often characterized, e.g., by multiple tissue antibodies, and/or a specific nonhistone ANA (SS-B)), diabetes mellitus (often characterized, e.g., by cell-mediated and humoral islet cell antibodies), and adrenergic drug resistance (including adrenergic drug resistance with asthma or cystic fibrosis) (often characterized, e.g., by beta-adrenergic receptor antibodies).

[0468] Additional disorders that may have an autoimmune component that may be treated, prevented, diagnosed and/or prognosed with the compositions of the invention include, but are not limited to, chronic active hepatitis (often characterized, e.g., by smooth muscle antibodies), primary biliary cirrhosis (often characterized, e.g., by mitochondria antibodies), other endocrine gland failure (often characterized, e.g., by specific tissue antibodies in some cases), vitiligo (often characterized, e.g., by melanocyte antibodies), vasculitis (often characterized, e.g., by Ig and complement in vessel walls and/or low serum complement), post-MI (often characterized, e.g., by myocardial antibodies), cardiomyopathy syndrome (often characterized, e.g., by myocardial antibodies), urticaria (often characterized, e.g., by IgG and IgM antibodies to IgE), atopic dermatitis (often characterized, e.g., by IgG and IgM antibodies to IgE), asthma (often characterized, e.g., by IgG and IgM antibodies to IgE), and many other inflammatory, granulomatous, degenerative, and atrophic disorders.

[0469] In a preferred embodiment, the autoimmune diseases and disorders and/or conditions associated with the diseases and disorders recited above are treated, prevented, diagnosed and/or prognosed using for example, antagonists or agonists, polypeptides or polynucleotides, or antibodies of the present invention. In a specific preferred embodiment, rheumatoid arthritis is treated, prevented, and/or diagnosed using polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention.

of self as foreign material by immune cells. This inappropriate recognition results in an immune response leading to the destruction of the host tissue. Therefore, the administration of polynucleotides and polypeptides of the invention that can inhibit an immune response, particularly the proliferation, differentiation, or chemotaxis of T-cells, may be an effective therapy in preventing autoimmune disorders.

[0465] Autoimmune diseases or disorders that may be treated, prevented, diagnosed and/or prognosed by polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention include, but are not limited to, one or more of the following: systemic lupus erythematosus, rheumatoid arthritis, ankylosing spondylitis, multiple sclerosis, autoimmune thyroiditis, Hashimoto's thyroiditis, autoimmune hemolytic anemia, hemolytic anemia, thrombocytopenia, autoimmune thrombocytopenia purpura, autoimmune neonatal thrombocytopenia, idiopathic thrombocytopenia purpura, purpura (e.g., Henloch-Scoenlein purpura), autoimmunocytopenia, Goodpasture's syndrome, Pemphigus vulgaris, myasthenia gravis, Grave's disease (hyperthyroidism), and insulin-resistant diabetes mellitus.

[0466] Additional disorders that are likely to have an autoimmune component that may be treated, prevented, and/or diagnosed with the compositions of the invention include, but are not limited to, type II collagen-induced arthritis, antiphospholipid syndrome, dermatitis, allergic encephalomyelitis, myocarditis, relapsing polychondritis, rheumatic heart disease, neuritis, uveitis ophthalmia, polyendocrinopathies, Reiter's Disease, Stiff-Man Syndrome, autoimmune pulmonary inflammation, autism, Guillain-Barre Syndrome, insulin dependent diabetes mellitus, and autoimmune inflammatory eye disorders.

[0467] Additional disorders that are likely to have an autoimmune component that may be treated, prevented, diagnosed and/or prognosed with the compositions of the invention include, but are not limited to, scleroderma with anti-collagen antibodies (often characterized, e.g., by nucleolar and other nuclear antibodies), mixed connective tissue disease (often characterized, e.g., by antibodies to extractable nuclear antigens (e.g., ribonucleoprotein)), polymyositis (often characterized, e.g., by nonhistone ANA), pernicious anemia (often characterized, e.g., by antiparietal cell, microsomes, and intrinsic factor antibodies), idiopathic Addison's disease (often characterized, e.g., by humoral and

pharyngeal pouch syndrome, 22q11.2 deletion, chronic mucocutaneous candidiasis, natural killer cell deficiency (NK), idiopathic CD4+ T-lymphocytopenia, immunodeficiency with predominant T cell defect (unspecified), and unspecified immunodeficiency of cell mediated immunity.

[0460] In specific embodiments, DiGeorge anomaly or conditions associated with DiGeorge anomaly are treated, prevented, diagnosed, and/or prognosed using polypeptides or polynucleotides of the invention, or antagonists or agonists thereof.

[0461] Other immunodeficiencies that may be treated, prevented, diagnosed, and/or prognosed using polypeptides or polynucleotides of the invention, and/or agonists or antagonists thereof, include, but are not limited to, chronic granulomatous disease, Chédiak-Higashi syndrome, myeloperoxidase deficiency, leukocyte glucose-6-phosphate dehydrogenase deficiency, X-linked lymphoproliferative syndrome (XLP), leukocyte adhesion deficiency, complement component deficiencies (including C1, C2, C3, C4, C5, C6, C7, C8 and/or C9 deficiencies), reticular dysgenesis, thymic aplasia-aplasia, immunodeficiency with thymoma, severe congenital leukopenia, dysplasia with immunodeficiency, neonatal neutropenia, short limbed dwarfism, and Nezelof syndrome-combined immunodeficiency with Igs.

[0462] In a preferred embodiment, the immunodeficiencies and/or conditions associated with the immunodeficiencies recited above are treated, prevented, diagnosed and/or prognosed using polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention.

[0463] In a preferred embodiment polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention could be used as an agent to boost immunoresponsiveness among immunodeficient individuals. In specific embodiments, polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention could be used as an agent to boost immunoresponsiveness among B cell and/or T cell immunodeficient individuals.

[0464] The polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in treating, preventing, diagnosing and/or prognosing autoimmune disorders. Many autoimmune disorders result from inappropriate recognition

diseases and disorders of the immune system and/or to inhibit or enhance an immune response generated by cells associated with the tissue(s) in which the polypeptide of the invention is expressed, including one, two, three, four, five, or more tissues disclosed in Table 1, column 8 (Tissue Distribution Library Code).

[0457] Polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in treating, preventing, diagnosing, and/or prognosing immunodeficiencies, including both congenital and acquired immunodeficiencies. Examples of B cell immunodeficiencies in which immunoglobulin levels B cell function and/or B cell numbers are decreased include: X-linked agammaglobulinemia (Bruton's disease), X-linked infantile agammaglobulinemia, X-linked immunodeficiency with hyper IgM, non X-linked immunodeficiency with hyper IgM, X-linked lymphoproliferative syndrome (XLP), agammaglobulinemia including congenital and acquired agammaglobulinemia, adult onset agammaglobulinemia, late-onset agammaglobulinemia, dysgammaglobulinemia, hypogammaglobulinemia, unspecified hypogammaglobulinemia, recessive agammaglobulinemia (Swiss type), Selective IgM deficiency, selective IgA deficiency, selective IgG subclass deficiencies, IgG subclass deficiency (with or without IgA deficiency), Ig deficiency with increased IgM, IgG and IgA deficiency with increased IgM, antibody deficiency with normal or elevated Igs, Ig heavy chain deletions, kappa chain deficiency, B cell lymphoproliferative disorder (BLPD), common variable immunodeficiency (CVID), common variable immunodeficiency (CVI) (acquired), and transient hypogammaglobulinemia of infancy.

[0458] In specific embodiments, ataxia-telangiectasia or conditions associated with ataxia-telangiectasia are treated, prevented, diagnosed, and/or prognosing using the polypeptides or polynucleotides of the invention, and/or agonists or antagonists thereof.

[0459] Examples of congenital immunodeficiencies in which T cell and/or B cell function and/or number is decreased include, but are not limited to: DiGeorge anomaly, severe combined immunodeficiencies (SCID) (including, but not limited to, X-linked SCID, autosomal recessive SCID, adenosine deaminase deficiency, purine nucleoside phosphorylase (PNP) deficiency, Class II MHC deficiency (Bare lymphocyte syndrome), Wiskott-Aldrich syndrome, and ataxia telangiectasia), thymic hypoplasia, third and fourth

composition comprising an amount of an isolated CD-like polypeptide, or antagonist thereof, effective to decrease the CD-like polypeptide activity level in such an individual.

[0452] In certain embodiments, a polypeptide of the invention, or polynucleotides, antibodies, agonists, or antagonists corresponding to that polypeptide, may be used to diagnose and/or prognose diseases and/or disorders associated with the tissue(s) in which the polypeptide of the invention is expressed, including one, two, three, four, five, or more tissues disclosed in Table 1, column 8 (Tissue Distribution Library Code).

[0453] Thus, polynucleotides, translation products and antibodies of the invention are useful in the diagnosis, prognosis, prevention, and/or treatment of diseases and/or disorders associated with activities that include, but are not limited to, blood coagulation, fibrinolysis, complement activation, extracellular matrix turnover, cell migration and prohormone activation.

[0454] More generally, polynucleotides, translation products and antibodies corresponding to this gene may be useful for the diagnosis, prognosis, prevention, and/or treatment of diseases and/or disorders associated with the following systems.

Immune Activity

[0455] Polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention may be useful in treating, preventing, diagnosing and/or prognosing diseases, disorders, and/or conditions of the immune system, by, for example, activating or inhibiting the proliferation, differentiation, or mobilization (chemotaxis) of immune cells. Immune cells develop through a process called hematopoiesis, producing myeloid (platelets, red blood cells, neutrophils, and macrophages) and lymphoid (B and T lymphocytes) cells from pluripotent stem cells. The etiology of these immune diseases, disorders, and/or conditions may be genetic, somatic, such as cancer and some autoimmune diseases, acquired (e.g., by chemotherapy or toxins), or infectious. Moreover, polynucleotides, polypeptides, antibodies, and/or agonists or antagonists of the present invention can be used as a marker or detector of a particular immune system disease or disorder.

[0456] In another embodiment, a polypeptide of the invention, or polynucleotides, antibodies, agonists, or antagonists corresponding to that polypeptide, may be used to treat

limited to, antimetabolites (e.g., methotrexate, 6-mercaptopurine, 6-thioguanine, cytarabine, 5-fluorouracil decarbazine), alkylating agents (e.g., mechlorethamine, thioepa chlorambucil, melphalan, carmustine (BSNU) and lomustine (CCNU), cyclophosphamide, busulfan, dibromomannitol, streptozotocin, mitomycin C, and cis- dichlorodiamine platinum (II) (DDP) cisplatin), anthracyclines (e.g., daunorubicin (formerly daunomycin) and doxorubicin), antibiotics (e.g., dactinomycin (formerly actinomycin), bleomycin, mithramycin, and anthramycin (AMC)), and anti-mitotic agents (e.g., vincristine and vinblastine).

[0449] By "cytotoxic prodrug" is meant a non-toxic compound that is converted by an enzyme, normally present in the cell, into a cytotoxic compound. Cytotoxic prodrugs that may be used according to the methods of the invention include, but are not limited to, glutamyl derivatives of benzoic acid mustard alkylating agent, phosphate derivatives of etoposide or mitomycin C, cytosine arabinoside, daunorubisin, and phenoxyacetamide derivatives of doxorubicin.

[0450] It will be appreciated that conditions caused by a decrease in the standard or normal level of CD-like polypeptide activity in an individual, particularly disorders of the immune system, can be treated by administration of CD-like polypeptides of the invention (e.g., in the form of soluble extracellular domain or cells expressing the complete protein) or agonist. Thus, the invention also provides a method of treatment of an individual in need of an increased level of CD-like polypeptide activity comprising administering to such an individual a pharmaceutical composition comprising an amount of an isolated CD-like polypeptides of the invention, or agonist thereof (e.g, an agonistic CD-like polypeptides antibody), effective to increase the CD-like polypeptides activity level in such an individual.

[0451] It will also be appreciated that conditions caused by a increase in the standard or normal level of CD-like polypeptide activity in an individual, particularly disorders of the immune system, can be treated by administration of CD-like polypeptides of the invention (e.g., in the form of soluble extracellular domain or cells expressing the complete protein) or antagonist (e.g, an antagonistic CD-like polypeptide antibody). Thus, the invention also provides a method of treatment of an individual in need of an decreased level of CD-like polypeptide activity comprising administering to such an individual a pharmaceutical

or ribozymes) or double stranded nucleic acid (e.g., DNA that can integrate into the cell's genome or replicate episomally and that can be transcribed) into the targeted cell.

[0446] In another embodiment, the invention provides a method for the specific destruction of cells (e.g., the destruction of tumor cells) by administering polypeptides of the invention (e.g., CD-like polypeptides or anti-CD-like polypeptides antibodies) in association with toxins or cytotoxic prodrugs.

[0447] By "toxin" is meant compounds that bind and activate endogenous cytotoxic effector systems, radioisotopes, holotoxins, modified toxins, catalytic subunits of toxins, cytotoxins (cytotoxic agents), or any molecules or enzymes not normally present in or on the surface of a cell that under defined conditions cause the cell's death. Toxins that may be used according to the methods of the invention include, but are not limited to, radioisotopes known in the art, compounds such as, for example, antibodies (or complement fixing containing portions thereof) that bind an inherent or induced endogenous cytotoxic effector system, thymidine kinase, endonuclease, RNase, alpha toxin, ricin, abrin, *Pseudomonas* exotoxin A, diphtheria toxin, saporin, momordin, gelonin, pokeweed antiviral protein, alpha-sarcin and cholera toxin. "Toxin" also includes a cytostatic or cytocidal agent, a therapeutic agent or a radioactive metal ion, e.g., alpha-emitters such as, for example, ^{213}Bi , or other radioisotopes such as, for example, ^{103}Pd , ^{133}Xe , ^{131}I , ^{68}Ge , ^{57}Co , ^{65}Zn , ^{85}Sr , ^{32}P , ^{35}S , ^{90}Y , ^{153}Sm , ^{153}Gd , ^{169}Yb , ^{51}Cr , ^{54}Mn , ^{75}Se , ^{113}Sn , $^{90}\text{Yttrium}$, ^{117}Tin , $^{186}\text{Rhenium}$, $^{166}\text{Holmium}$, and $^{188}\text{Rhenium}$; luminescent labels, such as luminol; and fluorescent labels, such as fluorescein and rhodamine, and biotin.

[0448] Techniques known in the art may be applied to label antibodies of the invention. Such techniques include, but are not limited to, the use of bifunctional conjugating agents (see e.g., U.S. Patent Nos. 5,756,065; 5,714,631; 5,696,239; 5,652,361; 5,505,931; 5,489,425; 5,435,990; 5,428,139; 5,342,604; 5,274,119; 4,994,560; and 5,808,003; the contents of each of which are hereby incorporated by reference in its entirety). A cytotoxin or cytotoxic agent includes any agent that is detrimental to cells. Examples include paclitaxol, cytochalasin B, gramicidin D, ethidium bromide, emetine, mitomycin, etoposide, tenoposide, vincristine, vinblastine, colchicin, doxorubicin, daunorubicin, dihydroxy anthracin dione, mitoxantrone, mithramycin, actinomycin D, 1-dehydrotestosterone, glucocorticoids, procaine, tetracaine, lidocaine, propranolol, and puromycin and analogs or homologs thereof. Therapeutic agents include, but are not

the diseases associated with the biological activity. Thus, the polynucleotides and polypeptides, and agonists or antagonists could be used to treat the associated disease.

[0443] The CD-like polypeptides of the invention may be involved in the immune and/or inflammatory response. For example, CD-like polypeptides of the invention (a) may be involved in cell-cell interactions, cell-surface recognition, intercellular communication; (b) act as cytokine receptors; (c) provide structural links between the extracellular environment and the cytoskeleton-signaling network and thus help to regulate cell migration, differentiation, cell cycle progression, apoptosis, phagocytosis, ECM assembly, and metalloproteinase activity; (d) may be involved in transmembrane signal transduction and regulation of cellular proliferation, development, motility, and tumor cell growth and metastasis, in a number of different cell types; (e) may bind a wide variety of pathogens and play a potential role in host defence; or (f) be involved in regulation of the complement activation system.

[0444] Thus, a CD-like polypeptides of the invention may be useful as a therapeutic molecule. It could be used to control the proliferation, activation, maturation, survival, and/or differentiation of hematopoietic cells, in particular B- and T-cells. Particularly, CD-like polypeptides of the invention may be a useful therapeutic to mediate immune modulation. This control of immune cells would be particularly important in the treatment, diagnosis, detection, and/or prevention of immune disorders, such as autoimmune diseases or immunosuppression (see below). Preferably, treatment, diagnosis, detection, and/or prevention of immune disorders could be carried out using a secreted form of a CD-like polypeptides of the invention, gene therapy, or ex vivo applications. Moreover, inhibitors of CD-like polypeptides of the invention, either blocking antibodies or mutant forms, could modulate the expression of CD-like polypeptides of the invention. These inhibitors

of the present invention. Suitable delivery vehicles for use with systemic administration comprise liposomes comprising ligands for targeting the vehicle to a particular site. In specific embodiments, suitable delivery vehicles for use with systemic administration comprise liposomes comprising polypeptides of the invention for targeting the vehicle to a particular site.

[0439] Preferred methods of systemic administration, include intravenous injection, aerosol, oral and percutaneous (topical) delivery. Intravenous injections can be performed using methods standard in the art. Aerosol delivery can also be performed using methods standard in the art (see, for example, Stribling et al., Proc. Natl. Acad. Sci. USA 189:11277-11281, 1992, which is incorporated herein by reference). Oral delivery can be performed by complexing a polynucleotide construct of the present invention to a carrier capable of withstanding degradation by digestive enzymes in the gut of an animal. Examples of such carriers, include plastic capsules or tablets, such as those known in the art. Topical delivery can be performed by mixing a polynucleotide construct of the present invention with a lipophilic reagent (e.g., DMSO) that is capable of passing into the skin.

[0440] Determining an effective amount of substance to be delivered can depend upon a number of factors including, for example, the chemical structure and biological activity of the substance, the age and weight of the animal, the precise condition requiring treatment and its severity, and the route of administration. The frequency of treatments depends upon a number of factors, such as the amount of polynucleotide constructs administered per dose, as well as the health and history of the subject. The precise amount, number of doses, and timing of doses will be determined by the attending physician or veterinarian.

[0441] Therapeutic compositions of the present invention can be administered to any animal, preferably to mammals and birds. Preferred mammals include humans, dogs, cats, mice, rats, rabbits sheep, cattle, horses and pigs, with humans being particularly preferred.

Biological Activities

[0442] Polynucleotides or polypeptides, or agonists or antagonists of the present invention, can be used in assays to test for one or more biological activities. If these polynucleotides or polypeptides, or agonists or antagonists of the present invention, do exhibit activity in a particular assay, it is likely that these molecules may be involved in

[0433] The promoter-targeting sequence construct is taken up by cells. Homologous recombination between the construct and the endogenous sequence takes place, such that an endogenous sequence is placed under the control of the promoter. The promoter then drives the expression of the endogenous sequence.

[0434] The polynucleotide encoding a polypeptide of the present invention may contain a secretory signal sequence that facilitates secretion of the protein. Typically, the signal sequence is positioned in the coding region of the polynucleotide to be expressed towards or at the 5' end of the coding region. The signal sequence may be homologous or heterologous to the polynucleotide of interest and may be homologous or heterologous to the cells to be transfected. Additionally, the signal sequence may be chemically synthesized using methods known in the art.

[0435] Any mode of administration of any of the above-described polynucleotides constructs can be used so long as the mode results in the expression of one or more molecules in an amount sufficient to provide a therapeutic effect. This includes direct needle injection, systemic injection, catheter infusion, biolistic injectors, particle accelerators (i.e., "gene guns"), gelfoam sponge depots, other commercially available depot materials, osmotic pumps (e.g., Alza minipumps), oral or suppositorial solid (tablet or pill) pharmaceutical formulations, and decanting or topical applications during surgery. For example, direct injection of naked calcium phosphate-precipitated plasmid into rat liver and rat spleen or a protein-coated plasmid into the portal vein has resulted in gene expression of the foreign gene in the rat livers (Kaneda et al., Science 243:375 (1989)).

[0436] A preferred method of local administration is by direct injection. Preferably, a recombinant molecule of the present invention complexed with a delivery vehicle is administered by direct injection into or locally within the area of arteries. Administration of a composition locally within the area of arteries refers to injecting the composition centimeters and preferably, millimeters within arteries.

[0437] Another method of local administration is to contact a polynucleotide construct of the present invention in or around a surgical wound. For example, a patient can undergo surgery and the polynucleotide construct can be coated on the surface of tissue inside the wound or the construct can be injected into areas of tissue inside the wound.

[0438] Therapeutic compositions useful in systemic administration, include recombinant molecules of the present invention complexed to a targeted delivery vehicle

or *in vivo*. The transduced cells will contain the polynucleotide construct integrated into its genome, and will express a polypeptide of the invention.

[0429] Another method of gene therapy involves operably associating heterologous control regions and endogenous polynucleotide sequences (e.g. encoding a polypeptide of the present invention) via homologous recombination (see, e.g., U.S. Patent No. 5,641,670, issued June 24, 1997; International Publication No. WO 96/29411, published September 26, 1996; International Publication No. WO 94/12650, published August 4, 1994; Koller et al., Proc. Natl. Acad. Sci. USA 86:8932-8935 (1989); and Zijlstra et al., Nature 342:435-438 (1989), which are herein incorporated by reference. This method involves the activation of a gene which is present in the target cells, but which is not normally expressed in the cells, or is expressed at a lower level than desired.

[0430] Polynucleotide constructs are made, using standard techniques known in the art, which contain the promoter with targeting sequences flanking the promoter. Suitable promoters are described herein. The targeting sequence is sufficiently complementary to an endogenous sequence to permit homologous recombination of the promoter-targeting sequence with the endogenous sequence. The targeting sequence will be sufficiently near the 5' end of the desired endogenous polynucleotide sequence so the promoter will be operably linked to the endogenous sequence upon homologous recombination.

[0431] The promoter and the targeting sequences can be amplified using PCR. Preferably, the amplified promoter contains distinct restriction enzyme sites on the 5' and 3' ends. Preferably, the 3' end of the first targeting sequence contains the same restriction enzyme site as the 5' end of the amplified promoter and the 5' end of the second targeting sequence contains the same restriction site as the 3' end of the amplified promoter. The amplified promoter and targeting sequences are digested and ligated together.

[0432] The promoter-targeting sequence construct is delivered to the cells, either as naked polynucleotide, or in conjunction with transfection-facilitating agents, such as liposomes, viral sequences, viral particles, whole viruses, lipofection, precipitating agents, etc., described in more detail above. The P promoter-targeting sequence can be delivered by any method, included direct needle injection, intravenous injection, topical administration, catheter infusion, particle accelerators, etc. The methods are described in more detail below.

the adenovirus vector Ad2 is useful and can be grown in human 293 cells. These cells contain the E1 region of adenovirus and constitutively express E1a and E1b, which complement the defective adenoviruses by providing the products of the genes deleted from the vector. In addition to Ad2, other varieties of adenovirus (e.g., Ad3, Ad5, and Ad7) are also useful in the present invention.

[0426] Preferably, the adenoviruses used in the present invention are replication deficient. Replication deficient adenoviruses require the aid of a helper virus and/or packaging cell line to form infectious particles. The resulting virus is capable of infecting cells and can express a polynucleotide of interest which is operably linked to a promoter, but cannot replicate in most cells. Replication deficient adenoviruses may be deleted in one or more of all or a portion of the following genes: E1a, E1b, E3, E4, E2a, or L1 through L5.

[0427] In certain other embodiments, the cells are engineered, *ex vivo* or *in vivo*, using an adeno-associated virus (AAV). AAVs are naturally occurring defective viruses that require helper viruses to produce infectious particles (Muzyczka, N., *Curr. Topics in Microbiol. Immunol.* 158:97 (1992)). It is also one of the few viruses that may integrate its DNA into non-dividing cells. Vectors containing as little as 300 base pairs of AAV can be packaged and can integrate, but space for exogenous DNA is limited to about 4.5 kb. Methods for producing and using such AAVs are known in the art. See, for example, U.S. Patent Nos. 5,139,941, 5,173,414, 5,354,678, 5,436,146, 5,474,935, 5,478,745, and 5,589,377.

[0428] For example, an appropriate AAV vector for use in the present invention will include all the sequences necessary for DNA replication, encapsidation, and host-cell integration. The polynucleotide construct is inserted into the AAV vector using standard cloning methods, such as those found in Sambrook et al., *Molecular Cloning: A Laboratory Manual*, Cold Spring Harbor Press (1989). The recombinant AAV vector is then transfected into packaging cells which are infected with a helper virus, using any standard technique, including lipofection, electroporation, calcium phosphate precipitation, etc. Appropriate helper viruses include adenoviruses, cytomegaloviruses, vaccinia viruses, or herpes viruses. Once the packaging cells are transfected and infected, they will produce infectious AAV viral particles which contain the polynucleotide construct. These viral particles are then used to transduce eukaryotic cells, either *ex vivo*

[0422] The retroviral plasmid vector is employed to transduce packaging cell lines to form producer cell lines. Examples of packaging cells which may be transfected include, but are not limited to, the PE501, PA317, R-2, R-AM, PA12, T19-14X, VT-19-17-H2, RCRE, RCRIP, GP+E-86, GP+envAm12, and DAN cell lines as described in Miller, Human Gene Therapy 1:5-14 (1990), which is incorporated herein by reference in its entirety. The vector may transduce the packaging cells through any means known in the art. Such means include, but are not limited to, electroporation, the use of liposomes, and CaPO_4 precipitation. In one alternative, the retroviral plasmid vector may be encapsulated into a liposome, or coupled to a lipid, and then administered to a host.

[0423] The producer cell line generates infectious retroviral vector particles which include polynucleotide encoding a polypeptide of the present invention. Such retroviral vector particles then may be employed, to transduce eukaryotic cells, either *in vitro* or *in vivo*. The transduced eukaryotic cells will express a polypeptide of the present invention.

[0424] In certain other embodiments, cells are engineered, *ex vivo* or *in vivo*, with polynucleotide contained in an adenovirus vector. Adenovirus can be manipulated such that it encodes and expresses a polypeptide of the present invention, and at the same time is inactivated in terms of its ability to replicate in a normal lytic viral life cycle. Adenovirus expression is achieved without integration of the viral DNA into the host cell chromosome, thereby alleviating concerns about insertional mutagenesis. Furthermore, adenoviruses have been used as live enteric vaccines for many years with an excellent safety profile (Schwartz et al. Am. Rev. Respir. Dis. 109:233-238 (1974)). Finally, adenovirus mediated gene transfer has been demonstrated in a number of instances including transfer of alpha-1-antitrypsin and CFTR to the lungs of cotton rats (Rosenfeld, M. A. et al. (1991) Science 252:431-434; Rosenfeld et al., (1992) Cell 68:143-155). Furthermore, extensive studies to attempt to establish adenovirus as a causative agent in human cancer were uniformly negative (Green, M. et al. (1979) Proc. Natl. Acad. Sci. USA 76:6606).

[0425] Suitable adenoviral vectors useful in the present invention are described, for example, in Kozarsky and Wilson, Curr. Opin. Genet. Devel. 3:499-503 (1993); Rosenfeld et al., Cell 68:143-155 (1992); Engelhardt et al., Human Genet. Ther. 4:759-769 (1993); Yang et al., Nature Genet. 7:362-369 (1994); Wilson et al., Nature 365:691-692 (1993); and U.S. Patent No. 5,652,224, which are herein incorporated by reference. For example,

buffer solution such as 10 mM Tris/NaCl, sonicated, and then the preformed liposomes are mixed directly with the DNA. The liposome and DNA form a very stable complex due to binding of the positively charged liposomes to the cationic DNA. SUVs find use with small nucleic acid fragments. LUVs are prepared by a number of methods, well known in the art. Commonly used methods include Ca^{2+} -EDTA chelation (Papahadjopoulos et al., Biochim. Biophys. Acta (1975) 394:483; Wilson et al., Cell 17:77 (1979)); ether injection (Deamer, D. and Bangham, A., Biochim. Biophys. Acta 443:629 (1976); Ostro et al., Biochem. Biophys. Res. Commun. 76:836 (1977); Fraley et al., Proc. Natl. Acad. Sci. USA 76:3348 (1979)); detergent dialysis (Enoch, H. and Strittmatter, P., Proc. Natl. Acad. Sci. USA 76:145 (1979)); and reverse-phase evaporation (REV) (Fraley et al., J. Biol. Chem. 255:10431 (1980); Szoka, F. and Papahadjopoulos, D., Proc. Natl. Acad. Sci. USA 75:145 (1978); Schaefer-Ridder et al., Science 215:166 (1982)), which are herein incorporated by reference.

[0419] Generally, the ratio of DNA to liposomes will be from about 10:1 to about 1:10. Preferably, the ratio will be from about 5:1 to about 1:5. More preferably, the ratio will be about 3:1 to about 1:3. Still more preferably, the ratio will be about 1:1.

[0420] U.S. Patent No. 5,676,954 (which is herein incorporated by reference) reports on the injection of genetic material, complexed with cationic liposomes carriers, into mice. U.S. Patent Nos. 4,897,355, 4,946,787, 5,049,386, 5,459,127, 5,589,466, 5,693,622, 5,580,859, 5,703,055, and international publication no. WO 94/9469 (which are herein incorporated by reference) provide cationic lipids for use in transfecting DNA into cells and mammals. U.S. Patent Nos. 5,589,466, 5,693,622, 5,580,859, 5,703,055, and international publication no. WO 94/9469 provide methods for delivering DNA-cationic lipid complexes to mammals.

[0421] In certain embodiments, cells are engineered, *ex vivo* or *in vivo*, using a retroviral particle containing RNA which comprises a sequence encoding a polypeptide of the present invention. Retroviruses from which the retroviral plasmid vectors may be derived include, but are not limited to, Moloney Murine Leukemia Virus, spleen necrosis virus, Rous sarcoma Virus, Harvey Sarcoma Virus, avian leukosis virus, gibbon ape leukemia virus, human immunodeficiency virus, Myeloproliferative Sarcoma Virus, and mammary tumor virus.

USA 84:7413-7417, which is herein incorporated by reference. Similar methods can be used to prepare liposomes from other cationic lipid materials.

[0416] Similarly, anionic and neutral liposomes are readily available, such as from Avanti Polar Lipids (Birmingham, Ala.), or can be easily prepared using readily available materials. Such materials include phosphatidyl, choline, cholesterol, phosphatidyl ethanolamine, dioleoylphosphatidyl choline (DOPC), dioleoylphosphatidyl glycerol (DOPG), dioleoylphosphatidyl ethanolamine (DOPE), among others. These materials can also be mixed with the DOTMA and DOTAP starting materials in appropriate ratios. Methods for making liposomes using these materials are well known in the art.

[0417] For example, commercially dioleoylphosphatidyl choline (DOPC), dioleoylphosphatidyl glycerol (DOPG), and dioleoylphosphatidyl ethanolamine (DOPE) can be used in various combinations to make conventional liposomes, with or without the addition of cholesterol. Thus, for example, DOPG/DOPC vesicles can be prepared by drying 50 mg each of DOPG and DOPC under a stream of nitrogen gas into a sonication vial. The sample is placed under a vacuum pump overnight and is hydrated the following day with deionized water. The sample is then sonicated for 2 hours in a capped vial, using a Heat Systems model 350 sonicator equipped with an inverted cup (bath type) probe at the maximum setting while the bath is circulated at 15EC. Alternatively, negatively charged vesicles can be prepared without sonication to produce multilamellar vesicles or by extrusion through nucleopore membranes to produce unilamellar vesicles of discrete size. Other methods are known and available to those of skill in the art.

[0418] The liposomes can comprise multilamellar vesicles (MLVs), small unilamellar vesicles (SUVs), or large unilamellar vesicles (LUVs), with SUVs being preferred. The various liposome-nucleic acid complexes are prepared using methods well known in the art. See, e.g., Straubinger et al., *Methods of Immunology* (1983), 101:512-527, which is herein incorporated by reference. For example, MLVs containing nucleic acid can be prepared by depositing a thin film of phospholipid on the walls of a glass tube and subsequently hydrating with a solution of the material to be encapsulated. SUVs are prepared by extended sonication of MLVs to produce a homogeneous population of unilamellar liposomes. The material to be entrapped is added to a suspension of preformed MLVs and then sonicated. When using liposomes containing cationic lipids, the dried lipid film is resuspended in an appropriate solution such as sterile water or an isotonic

tissues, throat or mucous membranes of the nose. In addition, naked DNA constructs can be delivered to arteries during angioplasty by the catheter used in the procedure.

[0411] The naked polynucleotides are delivered by any method known in the art, including, but not limited to, direct needle injection at the delivery site, intravenous injection, topical administration, catheter infusion, and so-called "gene guns". These delivery methods are known in the art.

[0412] The constructs may also be delivered with delivery vehicles such as viral sequences, viral particles, liposome formulations, lipofectin, precipitating agents, etc. Such methods of delivery are known in the art.

[0413] In certain embodiments, the polynucleotide constructs are complexed in a liposome preparation. Liposomal preparations for use in the instant invention include cationic (positively charged), anionic (negatively charged) and neutral preparations. However, cationic liposomes are particularly preferred because a tight charge complex can be formed between the cationic liposome and the polyanionic nucleic acid. Cationic liposomes have been shown to mediate intracellular delivery of plasmid DNA (Felgner et al., Proc. Natl. Acad. Sci. USA (1987) 84:7413-7416, which is herein incorporated by reference); mRNA (Malone et al., Proc. Natl. Acad. Sci. USA (1989) 86:6077-6081, which is herein incorporated by reference); and purified transcription factors (Debs et al., J. Biol. Chem. (1990) 265:10189-10192, which is herein incorporated by reference), in functional form.

[0414] Cationic liposomes are readily available. For example, N[1-2,3-dioleoyloxy)propyl]-N,N,N-triethylammonium (DOTMA) liposomes are particularly useful and are available under the trademark Lipofectin, from GIBCO BRL, Grand Island, N.Y. (See, also, Felgner et al., Proc. Natl Acad. Sci. USA (1987) 84:7413-7416, which is herein incorporated by reference). Other commercially available liposomes include transfectace (DDAB/DOPE) and DOTAP/DOPE (Boehringer).

[0415] Other cationic liposomes can be prepared from readily available materials using techniques well known in the art. See, e.g. PCT Publication No. WO 90/11092 (which is herein incorporated by reference) for a description of the synthesis of DOTAP (1,2-bis(oleoyloxy)-3-(trimethylammonio)propane) liposomes. Preparation of DOTMA liposomes is explained in the literature, see, e.g., P. Felgner et al., Proc. Natl. Acad. Sci.

[0407] Unlike other gene therapy techniques, one major advantage of introducing naked nucleic acid sequences into target cells is the transitory nature of the polynucleotide synthesis in the cells. Studies have shown that non-replicating DNA sequences can be introduced into cells to provide production of the desired polypeptide for periods of up to six months.

[0408] The polynucleotide construct can be delivered to the interstitial space of tissues within the an animal, including of muscle, skin, brain, lung, liver, spleen, bone marrow, thymus, heart, lymph, blood, bone, cartilage, pancreas, kidney, gall bladder, stomach, intestine, testis, ovary, uterus, rectum, nervous system, eye, gland, and connective tissue. Interstitial space of the tissues comprises the intercellular, fluid, mucopolysaccharide matrix among the reticular fibers of organ tissues, elastic fibers in the walls of vessels or chambers, collagen fibers of fibrous tissues, or that same matrix within connective tissue ensheathing muscle cells or in the lacunae of bone. It is similarly the space occupied by the plasma of the circulation and the lymph fluid of the lymphatic channels. Delivery to the interstitial space of muscle tissue is preferred for the reasons discussed below. They may be conveniently delivered by injection into the tissues comprising these cells. They are preferably delivered to and expressed in persistent, non-dividing cells which are differentiated, although delivery and expression may be achieved in non-differentiated or less completely differentiated cells, such as, for example, stem cells of blood or skin fibroblasts. *In vivo* muscle cells are particularly competent in their ability to take up and express polynucleotides.

[0409] For the naked nucleic acid sequence injection, an effective dosage amount of DNA or RNA will be in the range of from about 0.05 mg/kg body weight to about 50 mg/kg body weight. Preferably the dosage will be from about 0.005 mg/kg to about 20 mg/kg and more preferably from about 0.05 mg/kg to about 5 mg/kg. Of course, as the artisan of ordinary skill will appreciate, this dosage will vary according to the tissue site of injection. The appropriate and effective dosage of nucleic acid sequence can readily be determined by those of ordinary skill in the art and may depend on the condition being treated and the route of administration.

[0410] The preferred route of administration is by the parenteral route of injection into the interstitial space of tissues. However, other parenteral routes may also be used, such as, inhalation of an aerosol formulation particularly for delivery to lungs or bronchial

cells. The arterial cells may be reintroduced into the patient through direct injection to the artery, the tissues surrounding the artery, or through catheter injection.

[0403] As discussed in more detail below, the polynucleotide constructs can be delivered by any method that delivers injectable materials to the cells of an animal, such as, injection into the interstitial space of tissues (heart, muscle, skin, lung, liver, and the like). The polynucleotide constructs may be delivered in a pharmaceutically acceptable liquid or aqueous carrier.

[0404] In one embodiment, the polynucleotide of the present invention is delivered as a naked polynucleotide. The term "naked" polynucleotide, DNA or RNA refers to sequences that are free from any delivery vehicle that acts to assist, promote or facilitate entry into the cell, including viral sequences, viral particles, liposome formulations, lipofectin or precipitating agents and the like. However, the polynucleotide of the present invention can also be delivered in liposome formulations and lipofectin formulations and the like can be prepared by methods well known to those skilled in the art. Such methods are described, for example, in U.S. Patent Nos. 5,593,972, 5,589,466, and 5,580,859, which are herein incorporated by reference.

[0405] The polynucleotide vector constructs used in the gene therapy method are preferably constructs that will not integrate into the host genome nor will they contain sequences that allow for replication. Appropriate vectors include pWLNEO, pSV2CAT, pOG44, pXT1 and pSG available from Stratagene; pSVK3, pBPV, pMSG and pSVL available from Pharmacia; and pEF1/V5, pcDNA3.1, and pRc/CMV2 available from Invitrogen. Other suitable vectors will be readily apparent to the skilled artisan.

[0406] Any strong promoter known to those skilled in the art can be used for driving the expression of the polynucleotide sequence. Suitable promoters include adenoviral promoters, such as the adenoviral major late promoter; or heterologous promoters, such as the cytomegalovirus (CMV) promoter; the respiratory syncytial virus (RSV) promoter; inducible promoters, such as the MMT promoter, the metallothionein promoter; heat shock promoters; the albumin promoter; the ApoAI promoter; human globin promoters; viral thymidine kinase promoters, such as the Herpes Simplex thymidine kinase promoter; retroviral LTRs; the b-actin promoter; and human growth hormone promoters. The promoter also may be the native promoter for the polynucleotide of the present invention.

ng to about 10 ug, and preferably about 100 ng to about 1 ug, is sufficient to immobilize an adequate amount of binding agent.

[0400] Covalent attachment of binding agent to a solid support may generally be achieved by first reacting the support with a bifunctional reagent that will react with both the support and a functional group, such as a hydroxyl or amino group, on the binding agent. For example, the binding agent may be covalently attached to supports having an appropriate polymer coating using benzoquinone or by condensation of an aldehyde group on the support with an amine and an active hydrogen on the binding partner (see, e.g., Pierce Immunotechnology Catalog and Handbook, 1991, at A12-A13).

Gene Therapy Methods

[0401] Also encompassed by the invention are gene therapy methods for treating or preventing disorders, diseases and conditions. The gene therapy methods relate to the introduction of nucleic acid (DNA, RNA and antisense DNA or RNA) sequences into an animal to achieve expression of the polypeptide of the present invention. This method requires a polynucleotide which codes for a polypeptide of the present invention operatively linked to a promoter and any other genetic elements necessary for the expression of the polypeptide by the target tissue. Such gene therapy and delivery techniques are known in the art, see, for example, WO90/11092, which is herein incorporated by reference.

[0402] Thus, for example, cells from a patient may be engineered with a polynucleotide (DNA or RNA) comprising a promoter operably linked to a polynucleotide of the present invention ex vivo, with the engineered cells then being provided to a patient to be treated with the polypeptide of the present invention. Such methods are well-known in the art. For example, see Belldgrun, A., et al., J. Natl. Cancer Inst. 85: 207-216 (1993); Ferrantini, M. et al., Cancer Research 53: 1107-1112 (1993); Ferrantini, M. et al., J. Immunology 153: 4604-4615 (1994); Kaido, T., et al., Int. J. Cancer 60: 221-229 (1995); Ogura, H., et al., Cancer Research 50: 5102-5106 (1990); Santodonato, L., et al., Human Gene Therapy 7:1-10 (1996); Santodonato, L., et al., Gene Therapy 4:1246-1255 (1997); and Zhang, J.-F. et al., Cancer Gene Therapy 3: 31-38 (1996)), which are herein incorporated by reference. In one embodiment, the cells which are engineered are arterial

[0398] In a preferred embodiment, the assay involves the use of a binding agent(s) immobilized on a solid support to bind to and remove the polypeptide of the invention from the remainder of the sample. The bound polypeptide may then be detected using a detection reagent that contains a reporter group and specifically binds to the binding agent/polypeptide complex. Such detection reagents may comprise, for example, a binding agent that specifically binds to the polypeptide or an antibody or other agent that specifically binds to the binding agent, such as an anti-immunoglobulin, protein G, protein A or a lectin. Alternatively, a competitive assay may be utilized, in which a polypeptide is labeled with a reporter group and allowed to bind to the immobilized binding agent after incubation of the binding agent with the sample. The extent to which components of the sample inhibit the binding of the labeled polypeptide to the binding agent is indicative of the reactivity of the sample with the immobilized binding agent. Suitable polypeptides for use within such assays include polypeptides of the invention and portions thereof, or antibodies, to which the binding agent binds, as described above.

[0399] The solid support may be any material known to those of skill in the art to which polypeptides of the invention may be attached. For example, the solid support may be a test well in a microtiter plate or a nitrocellulose or other suitable membrane. Alternatively, the support may be a bead or disc, such as glass fiberglass, latex or a plastic material such as polystyrene or polyvinylchloride. The support may also be a magnetic particle or a fiber optic sensor, such as those disclosed, for example, in U.S. Patent No. 5,359,681. The binding agent may be immobilized on the solid support using a variety of techniques known to those of skill in the art, which are amply described in the patent and scientific literature. In the context of the present invention, the term "immobilization" refers to both noncovalent association, such as adsorption, and covalent attachment (which may be a direct linkage between the agent and functional groups on the support or may be a linkage by way of a cross-linking agent). Immobilization by adsorption to a well in a microtiter plate or to a membrane is preferred. In such cases, adsorption may be achieved by contacting the binding agent, in a suitable buffer, with the solid support for the suitable amount of time. The contact time varies with temperature, but is typically between about 1 hour and about 1 day. In general, contacting a well of plastic microtiter plate (such as polystyrene or polyvinylchloride) with an amount of binding agent ranging from about 10

chemiluminescent compound. The presence of the chemiluminescent-tagged antibody is then determined by detecting the presence of luminescence that arises during the course of a chemical reaction. Examples of particularly useful chemiluminescent labeling compounds are luminol, isoluminol, therrromatic acridinium ester, imidazole, acridinium salt and oxalate ester.

[0395] Likewise, a bioluminescent compound may be used to label the antibody of the present invention. Bioluminescence is a type of chemiluminescence found in biological systems in, which a catalytic protein increases the efficiency of the chemiluminescent reaction. The presence of a bioluminescent protein is determined by detecting the presence of luminescence. Important bioluminescent compounds for purposes of labeling are luciferin, luciferase and aequorin.

Methods for Detecting Diseases, Including Cancer

[0396] In general, a disease may be detected in a patient based on the presence of one or more proteins of the invention and/or polynucleotides encoding such proteins in a biological sample (for example, blood, sera, urine, and/or tumor biopsies) obtained from the patient. In other words, such proteins may be used as markers to indicate the presence or absence of a disease or disorder, including cancer and/or as described elsewhere herein. In addition, such proteins may be useful for the detection of other diseases and cancers. The binding agents provided herein generally permit detection of the level of antigen that binds to the agent in the biological sample. Polynucleotide primers and probes may be used to detect the level of mRNA encoding polypeptides of the invention, which is also indicative of the presence or absence of a disease or disorder, including cancer. In general, polypeptides of the invention should be present at a level that is at least three fold higher in diseased tissue than in normal tissue.

[0397] There are a variety of assay formats known to those of ordinary skill in the art for using a binding agent to detect polypeptide markers in a sample. See, e.g., Harlow and Lane, *supra*. In general, the presence or absence of a disease in a patient may be determined by (a) contacting a biological sample obtained from a patient with a binding agent; (b) detecting in the sample a level of polypeptide that binds to the binding agent; and (c) comparing the level of polypeptide with a predetermined cut-off value.

1980, Enzyme Immunoassay, CRC Press, Boca Raton, FL.; Ishikawa, E. et al., (eds.), 1981, Enzyme Immunoassay, Kigaku Shoin, Tokyo). The reporter enzyme which is bound to the antibody will react with an appropriate substrate, preferably a chromogenic substrate, in such a manner as to produce a chemical moiety which can be detected, for example, by spectrophotometric, fluorimetric or by visual means. Reporter enzymes which can be used to detectably label the antibody include, but are not limited to, malate dehydrogenase, staphylococcal nuclease, delta-5-steroid isomerase, yeast alcohol dehydrogenase, alpha-glycerophosphate dehydrogenase, triose phosphate isomerase, horseradish peroxidase, alkaline phosphatase, asparaginase, glucose oxidase, beta-galactosidase, ribonuclease, urease, catalase, glucose-6-phosphate dehydrogenase, glucoamylase and acetylcholinesterase. Additionally, the detection can be accomplished by colorimetric methods which employ a chromogenic substrate for the reporter enzyme. Detection may also be accomplished by visual comparison of the extent of enzymatic reaction of a substrate in comparison with similarly prepared standards.

[0391] Detection may also be accomplished using any of a variety of other immunoassays. For example, by radioactively labeling the antibodies or antibody fragments, it is possible to detect polypeptides through the use of a radioimmunoassay (RIA) (see, for example, Weintraub, B., Principles of Radioimmunoassays, Seventh Training Course on Radioligand Assay Techniques, The Endocrine Society, March, 1986, which is incorporated by reference herein). The radioactive isotope can be detected by means including, but not limited to, a gamma counter, a scintillation counter, or autoradiography.

[0392] It is also possible to label the antibody with a fluorescent compound. When the fluorescently labeled antibody is exposed to light of the proper wave length, its presence can then be detected due to fluorescence. Among the most commonly used fluorescent labeling compounds are fluorescein isothiocyanate, rhodamine, phycoerythrin, phycocyanin, allophycocyanin, ophthaldehyde and fluorescamine.

[0393] The antibody can also be detectably labeled using fluorescence emitting metals such as ^{152}Eu , or others of the lanthanide series. These metals can be attached to the antibody using such metal chelating groups as diethylenetriaminepentacetic acid (DTPA) or ethylenediaminetetraacetic acid (EDTA).

[0394] The antibody also can be detectably labeled by coupling it to a

monoclonal antibodies. Such antibodies can be produced using techniques described herein or otherwise known in the art. For example methods for producing chimeric antibodies are known in the art. See, for review, Morrison, *Science* 229:1202 (1985); Oi et al., *BioTechniques* 4:214 (1986); Cabilly et al., U.S. Patent No. 4,816,567; Taniguchi et al., EP 171496; Morrison et al., EP 173494; Neuberger et al., WO 8601533; Robinson et al., WO 8702671; Boulianne *et al.*, *Nature* 312:643 (1984); Neuberger *et al.*, *Nature* 314:268 (1985).

[0388] Additionally, any polypeptides of the invention whose presence can be detected, can be administered. For example, polypeptides of the invention labeled with a radio-opaque or other appropriate compound can be administered and visualized *in vivo*, as discussed, above for labeled antibodies. Further, such polypeptides can be utilized for *in vitro* diagnostic procedures.

[0389] A polypeptide-specific antibody or antibody fragment which has been labeled with an appropriate detectable imaging moiety, such as a radioisotope (for example, ^{131}I , ^{112}In , $^{99\text{m}}\text{Tc}$), a radio-opaque substance, or a material detectable by nuclear magnetic resonance, is introduced (for example, parenterally, subcutaneously or intraperitoneally) into the mammal to be examined for a disorder. It will be understood in the art that the size of the subject and the imaging system used will determine the quantity of imaging moiety needed to produce diagnostic images. In the case of a radioisotope moiety, for a human subject, the quantity of radioactivity injected will normally range from about 5 to 20 millicuries of $^{99\text{m}}\text{Tc}$. The labeled antibody or antibody fragment will then preferentially accumulate at the location of cells which contain the antigenic protein. *In vivo* tumor imaging is described in S.W. Burchiel et al., "Immunopharmacokinetics of Radiolabeled Antibodies and Their Fragments" (Chapter 13 in *Tumor Imaging: The Radiochemical Detection of Cancer*, S.W. Burchiel and B. A. Rhodes, eds., Masson Publishing Inc. (1982)).

[0390] With respect to antibodies, one of the ways in which an antibody of the present invention can be detectably labeled is by linking the same to a reporter enzyme and using the linked product in an enzyme immunoassay (EIA) (Voller, A., "The Enzyme Linked Immunosorbent Assay (ELISA)", 1978, Diagnostic Horizons 2:1-7, Microbiological Associates Quarterly Publication, Walkersville, MD); Voller et al., *J. Clin. Pathol.* 31:507-520 (1978); Butler, J.E., *Meth. Enzymol.* 73:482-523 (1981); Maggio, E. (ed.),

polypropylene, polyethylene, dextran, nylon, amylases, natural and modified celluloses, polyacrylamides, gabbros, and magnetite. The nature of the carrier can be either soluble to some extent or insoluble for the purposes of the present invention. The support material may have virtually any possible structural configuration so long as the coupled molecule is capable of binding to an antigen or antibody. Thus, the support configuration may be spherical, as in a bead, or cylindrical, as in the inside surface of a test tube, or the external surface of a rod. Alternatively, the surface may be flat such as a sheet, test strip, etc. Preferred supports include polystyrene beads. Those skilled in the art will know many other suitable carriers for binding antibody or antigen, or will be able to ascertain the same by use of routine experimentation.

[0385] The binding activity of a given lot of antibody or antigen polypeptide may be determined according to well known methods. Those skilled in the art will be able to determine operative and optimal assay conditions for each determination by employing routine experimentation.

[0386] In addition to assaying polypeptide levels or polynucleotide levels in a biological sample obtained from an individual, polypeptide or polynucleotide can also be detected *in vivo* by imaging. For example, in one embodiment of the invention, polypeptides and/or antibodies of the invention are used to image diseased cells, such as neoplasms. In another embodiment, polynucleotides of the invention (e.g., polynucleotides complementary to all or a portion of an mRNA) and/or antibodies (e.g., antibodies directed to any one or a combination of the epitopes of a polypeptide of the invention, antibodies directed to a conformational epitope of a polypeptide of the invention, or antibodies directed to the full length polypeptide expressed on the cell surface of a mammalian cell) are used to image diseased or neoplastic cells.

[0387] Antibody labels or markers for *in vivo* imaging of polypeptides of the invention include those detectable by X-radiography, NMR, MRI, CAT-scans or ESR. For X-radiography, suitable labels include radioisotopes such as barium or cesium, which emit detectable radiation but are not overtly harmful to the subject. Suitable markers for NMR and ESR include those with a detectable characteristic spin, such as deuterium, which may be incorporated into the antibody by labeling of nutrients for the relevant hybridoma. Where *in vivo* imaging is used to detect enhanced levels of polypeptides for diagnosis in humans, it may be preferable to use human antibodies or "humanized" chimeric

or peptide fragments thereof. This can be accomplished, for example, by immunofluorescence techniques employing a fluorescently labeled antibody coupled with light microscopic, flow cytometric, or fluorimetric detection.

[0381] The antibodies (or fragments thereof), and/or polypeptides of the present invention may, additionally, be employed histologically, as in immunofluorescence, immunoelectron microscopy or non-immunological assays, for in situ detection of gene products or conserved variants or peptide fragments thereof. In situ detection may be accomplished by removing a histological specimen from a patient, and applying thereto a labeled antibody or polypeptide of the present invention. The antibody (or fragment thereof) or polypeptide is preferably applied by overlaying the labeled antibody (or fragment) onto a biological sample. Through the use of such a procedure, it is possible to determine not only the presence of the gene product, or conserved variants or peptide fragments, or polypeptide binding, but also its distribution in the examined tissue. Using the present invention, those of ordinary skill will readily perceive that any of a wide variety of histological methods (such as staining procedures) can be modified in order to achieve such in situ detection.

[0382] Immunoassays and non-immunoassays for gene products or conserved variants or peptide fragments thereof will typically comprise incubating a sample, such as a biological fluid, a tissue extract, freshly harvested cells, or lysates of cells which have been incubated in cell culture, in the presence of a detectably labeled antibody capable of binding gene products or conserved variants or peptide fragments thereof, and detecting the bound antibody by any of a number of techniques well-known in the art.

[0383] The biological sample may be brought in contact with and immobilized onto a solid phase support or carrier such as nitrocellulose, or other solid support which is capable of immobilizing cells, cell particles or soluble proteins. The support may then be washed with suitable buffers followed by treatment with the detectably labeled antibody or detectable polypeptide of the invention. The solid phase support may then be washed with the buffer a second time to remove unbound antibody or polypeptide. Optionally the antibody is subsequently labeled. The amount of bound label on solid support may then be detected by conventional means.

[0384] By "solid phase support or carrier" is intended any support capable of binding an antigen or an antibody. Well-known supports or carriers include glass, polystyrene,

with classical immunohistological methods (Jalkanen et al., J. Cell. Biol. 101:976-985 (1985); Jalkanen, M., et al., J. Cell. Biol. 105:3087-3096 (1987)). Other antibody-based methods useful for detecting polypeptide gene expression include immunoassays, such as the enzyme linked immunosorbent assay (ELISA) and the radioimmunoassay (RIA). Suitable antibody assay labels are known in the art and include enzyme labels, such as, glucose oxidase, and radioisotopes, such as iodine (^{125}I , ^{121}I), carbon (^{14}C), sulfur (^{35}S), tritium (^3H), indium (^{112}In), and technetium ($^{99\text{m}}\text{Tc}$), and fluorescent labels, such as fluorescein and rhodamine, and biotin.

[0377] The tissue or cell type to be analyzed will generally include those which are known, or suspected, to express the gene of interest (such as, for example, cancer). The protein isolation methods employed herein may, for example, be such as those described in Harlow and Lane (Harlow, E. and Lane, D., 1988, "Antibodies: A Laboratory Manual", Cold Spring Harbor Laboratory Press, Cold Spring Harbor, New York), which is incorporated herein by reference in its entirety. The isolated cells can be derived from cell culture or from a patient. The analysis of cells taken from culture may be a necessary step in the assessment of cells that could be used as part of a cell-based gene therapy technique or, alternatively, to test the effect of compounds on the expression of the gene.

[0378] For example, antibodies, or fragments of antibodies, such as those described herein, may be used to quantitatively or qualitatively detect the presence of gene products or conserved variants or peptide fragments thereof. This can be accomplished, for example, by immunofluorescence techniques employing a fluorescently labeled antibody coupled with light microscopic, flow cytometric, or fluorimetric detection.

[0379] In a preferred embodiment, antibodies, or fragments of antibodies directed to any one or all of the predicted epitope domains of the polypeptides of the invention (shown in column 7 of Table 1) may be used to quantitatively or qualitatively detect the presence of gene products or conserved variants or peptide fragments thereof. This can be accomplished, for example, by immunofluorescence techniques employing a fluorescently labeled antibody coupled with light microscopic, flow cytometric, or fluorimetric detection.

[0380] In an additional preferred embodiment, antibodies, or fragments of antibodies directed to a conformational epitope of a polypeptide of the invention may be used to quantitatively or qualitatively detect the presence of gene products or conserved variants

from a second biological sample obtained from an individual not having the disorder or being determined by averaging levels from a population of individuals not having the disorder. As will be appreciated in the art, once a standard polypeptide level or mRNA level is known, it can be used repeatedly as a standard for comparison.

[0373] By "biological sample" is intended any biological sample obtained from an individual, cell line, tissue culture, or other source containing polypeptides of the invention (including portions thereof) or mRNA. As indicated, biological samples include body fluids (such as sera, plasma, urine, synovial fluid and spinal fluid) and tissue sources found to express the full length or fragments thereof of a polypeptide or mRNA. Methods for obtaining tissue biopsies and body fluids from mammals are well known in the art. Where the biological sample is to include mRNA, a tissue biopsy is the preferred source.

[0374] Total cellular RNA can be isolated from a biological sample using any suitable technique such as the single-step guanidinium-thiocyanate-phenol-chloroform method described in Chomczynski and Sacchi, *Anal. Biochem.* 162:156-159 (1987). Levels of mRNA encoding the polypeptides of the invention are then assayed using any appropriate method. These include Northern blot analysis, S1 nuclease mapping, the polymerase chain reaction (PCR), reverse transcription in combination with the polymerase chain reaction (RT-PCR), and reverse transcription in combination with the ligase chain reaction (RT-LCR).

[0375] The present invention also relates to diagnostic assays such as quantitative and diagnostic assays for detecting levels of polypeptides of the invention, in a biological sample (e.g., cells and tissues), including determination of normal and abnormal levels of polypeptides. Thus, for instance, a diagnostic assay in accordance with the invention for detecting over-expression of polypeptides of the invention compared to normal control tissue samples may be used to detect the presence of tumors. Assay techniques that can be used to determine levels of a polypeptide, such as a polypeptide of the present invention in a sample derived from a host are well-known to those of skill in the art. Such assay methods include radioimmunoassays, competitive-binding assays, Western Blot analysis and ELISA assays. Assaying polypeptide levels in a biological sample can occur using any art-known method.

[0376] Assaying polypeptide levels in a biological sample can occur using antibody-based techniques. For example, polypeptide expression in tissues can be studied

[0368] The compounds of the present invention are useful for diagnosis, treatment, prevention and/or prognosis of various disorders in mammals, preferably humans. Such disorders include, but are not limited to, those described herein under the section heading "Biological Activities".

[0369] For a number of disorders, substantially altered (increased or decreased) levels of gene expression can be detected in tissues, cells or bodily fluids (e.g., sera, plasma, urine, semen, synovial fluid or spinal fluid) taken from an individual having such a disorder, relative to a "standard" gene expression level, that is, the expression level in tissues or bodily fluids from an individual not having the disorder. Thus, the invention provides a diagnostic method useful during diagnosis of a disorder, which involves measuring the expression level of the gene encoding the polypeptide in tissues, cells or body fluid from an individual and comparing the measured gene expression level with a standard gene expression level, whereby an increase or decrease in the gene expression level(s) compared to the standard is indicative of a disorder. These diagnostic assays may be performed *in vivo* or *in vitro*, such as, for example, on blood samples, biopsy tissue or autopsy tissue.

[0370] The present invention is also useful as a prognostic indicator, whereby patients exhibiting enhanced or depressed gene expression will experience a worse clinical outcome relative to patients expressing the gene at a level nearer the standard level.

[0371] In certain embodiments, a polypeptide of the invention, or polynucleotides, antibodies, agonists, or antagonists corresponding to that polypeptide, may be used to diagnose and/or prognose diseases and/or disorders associated with the tissue(s) in which the polypeptide of the invention is expressed, including one, two, three, four, five, or more tissues disclosed in Table 1, column 8 (Tissue Distribution Library Code).

[0372] By "assaying the expression level of the gene encoding the polypeptide" is intended qualitatively or quantitatively measuring or estimating the level of the polypeptide of the invention or the level of the mRNA encoding the polypeptide of the invention in a first biological sample either directly (e.g., by determining or estimating absolute protein level or mRNA level) or relatively (e.g., by comparing to the polypeptide level or mRNA level in a second biological sample). Preferably, the polypeptide expression level or mRNA level in the first biological sample is measured or estimated and compared to a standard polypeptide level or mRNA level, the standard being taken

the disease, or may provide a means for detecting the disease prior to the appearance of actual clinical symptoms. A more definitive diagnosis of this type may allow health professionals to employ preventative measures or aggressive treatment earlier thereby preventing the development or further progression of the cancer.

[0365] Moreover, polypeptides of the present invention can be used to treat or prevent diseases or conditions such as, for example, neural disorders, immune system disorders, muscular disorders, reproductive disorders, gastrointestinal disorders, pulmonary disorders, cardiovascular disorders, renal disorders, proliferative disorders, and/or cancerous diseases and conditions. For example, patients can be administered a polypeptide of the present invention in an effort to replace absent or decreased levels of the polypeptide (e.g., insulin), to supplement absent or decreased levels of a different polypeptide (e.g., hemoglobin S for hemoglobin B, SOD, catalase, DNA repair proteins), to inhibit the activity of a polypeptide (e.g., an oncogene or tumor suppressor), to activate the activity of a polypeptide (e.g., by binding to a receptor), to reduce the activity of a membrane bound receptor by competing with it for free ligand (e.g., soluble TNF receptors used in reducing inflammation), or to bring about a desired response (e.g., blood vessel growth inhibition, enhancement of the immune response to proliferative cells or tissues).

[0366] Similarly, antibodies directed to a polypeptide of the present invention can also be used to treat disease (as described *supra*, and elsewhere herein). For example, administration of an antibody directed to a polypeptide of the present invention can bind, and/or neutralize the polypeptide, and/or reduce overproduction of the polypeptide. Similarly, administration of an antibody can activate the polypeptide, such as by binding to a polypeptide bound to a membrane (receptor).

[0367] At the very least, the polypeptides of the present invention can be used as molecular weight markers on SDS-PAGE gels or on molecular sieve gel filtration columns using methods well known to those of skill in the art. Polypeptides can also be used to raise antibodies, which in turn are used to measure protein expression from a recombinant cell, as a way of assessing transformation of the host cell. Moreover, the polypeptides of the present invention can be used to test the biological activities described herein.

Diagnostic Assays

art, compounds such as, for example, antibodies (or complement fixing containing portions thereof) that bind an inherent or induced endogenous cytotoxic effector system, thymidine kinase, endonuclease, RNase, alpha toxin, ricin, abrin, *Pseudomonas* exotoxin A, diphtheria toxin, saporin, momordin, gelonin, pokeweed antiviral protein, alpha-sarcin and cholera toxin. "Toxin" also includes a cytostatic or cytotoxic agent, a therapeutic agent or a radioactive metal ion, e.g., alpha-emitters such as, for example, ^{213}Bi , or other radioisotopes such as, for example, ^{103}Pd , ^{133}Xe , ^{131}I , ^{68}Ge , ^{57}Co , ^{65}Zn , ^{85}Sr , ^{32}P , ^{35}S , ^{90}Y , ^{153}Sm , ^{153}Gd , ^{169}Yb , ^{51}Cr , ^{54}Mn , ^{75}Se , ^{113}Sn , $^{90}\text{Yttrium}$, ^{117}Tin , $^{186}\text{Rhenium}$, $^{166}\text{Holmium}$, and $^{188}\text{Rhenium}$; luminescent labels, such as luminol; and fluorescent labels, such as fluorescein and rhodamine, and biotin. In a specific embodiment, the invention provides a method for the specific destruction of cells (e.g., the destruction of tumor cells) by administering polypeptides of the invention or antibodies of the invention in association with the radioisotope ^{90}Y . In another specific embodiment, the invention provides a method for the specific destruction of cells (e.g., the destruction of tumor cells) by administering polypeptides of the invention or antibodies of the invention in association with the radioisotope ^{111}In . In a further specific embodiment, the invention provides a method for the specific destruction of cells (e.g., the destruction of tumor cells) by administering polypeptides of the invention or antibodies of the invention in association with the radioisotope ^{131}I .

[0363] Techniques known in the art may be applied to label polypeptides of the invention (including antibodies). Such techniques include, but are not limited to, the use of bifunctional conjugating agents (see e.g., U.S. Patent Nos. 5,756,065; 5,714,631; 5,696,239; 5,652,361; 5,505,931; 5,489,425; 5,435,990; 5,428,139; 5,342,604; 5,274,119; 4,994,560; and 5,808,003; the contents of each of which are hereby incorporated by reference in its entirety).

[0364] Thus, the invention provides a diagnostic method of a disorder, which involves (a) assaying the expression level of a polypeptide of the present invention in cells or body fluid of an individual; and (b) comparing the assayed polypeptide expression level with a standard polypeptide expression level, whereby an increase or decrease in the assayed polypeptide expression level compared to the standard expression level is indicative of a disorder. With respect to cancer, the presence of a relatively high amount of transcript in biopsied tissue from an individual may indicate a predisposition for the development of

^{112}In , ^{111}In), and technetium (^{99}Tc , $^{99\text{m}}\text{Tc}$), thallium (^{201}Tl), gallium (^{68}Ga , ^{67}Ga), palladium (^{103}Pd), molybdenum (^{99}Mo), xenon (^{133}Xe), fluorine (^{18}F , ^{153}Sm , ^{177}Lu , ^{159}Gd , ^{149}Pm , ^{140}La , ^{175}Yb , ^{166}Ho , ^{90}Y , ^{47}Sc , ^{186}Re , ^{188}Re , ^{142}Pr , ^{105}Rh , ^{97}Ru), a radio-opaque substance, or a material detectable by nuclear magnetic resonance, is introduced (for example, parenterally, subcutaneously or intraperitoneally) into the mammal to be examined for immune system disorder. It will be understood in the art that the size of the subject and the imaging system used will determine the quantity of imaging moiety needed to produce diagnostic images. In the case of a radioisotope moiety, for a human subject, the quantity of radioactivity injected will normally range from about 5 to 20 millicuries of $^{99\text{m}}\text{Tc}$. The labeled antibody or antibody fragment will then preferentially accumulate at the location of cells which express the polypeptide encoded by a polynucleotide of the invention. *In vivo* tumor imaging is described in S.W. Burchiel et al., "Immunopharmacokinetics of Radiolabeled Antibodies and Their Fragments" (Chapter 13 in *Tumor Imaging: The Radiochemical Detection of Cancer*, S.W. Burchiel and B. A. Rhodes, eds., Masson Publishing Inc. (1982)).

[0360] In one embodiment, the invention provides a method for the specific delivery of compositions of the invention to cells by administering polypeptides of the invention (e.g., polypeptides encoded by polynucleotides of the invention and/or antibodies) that are associated with heterologous polypeptides or nucleic acids. In one example, the invention provides a method for delivering a therapeutic protein into the targeted cell. In another example, the invention provides a method for delivering a single stranded nucleic acid (e.g., antisense or ribozymes) or double stranded nucleic acid (e.g., DNA that can integrate into the cell's genome or replicate episomally and that can be transcribed) into the targeted cell.

[0361] In another embodiment, the invention provides a method for the specific destruction of cells (e.g., the destruction of tumor cells) by administering polypeptides of the invention in association with toxins or cytotoxic prodrugs.

[0362] By "toxin" is meant one or more compounds that bind and activate endogenous cytotoxic effector systems, radioisotopes, holotoxins, modified toxins, catalytic subunits of toxins, or any molecules or enzymes not normally present in or on the surface of a cell that under defined conditions cause the cell's death. Toxins that may be used according to the methods of the invention include, but are not limited to, radioisotopes known in the

Uses of the P lypeptides

[0355] Each of the polypeptides identified herein can be used in numerous ways. The following description should be considered exemplary and utilizes known techniques.

[0356] Polypeptides and antibodies directed to polypeptides of the present invention are useful to provide immunological probes for differential identification of the tissue(s) (e.g., immunohistochemistry assays such as, for example, ABC immunoperoxidase (Hsu et al., J. Histochem. Cytochem. 29:577-580 (1981)) or cell type(s) (e.g., immunocytochemistry assays).

[0357] Antibodies can be used to assay levels of polypeptides encoded by polynucleotides of the invention in a biological sample using classical immunohistological methods known to those of skill in the art (e.g., see Jalkanen, et al., J. Cell. Biol. 101:976-985 (1985); Jalkanen, et al., J. Cell. Biol. 105:3087-3096 (1987)). Other antibody-based methods useful for detecting protein gene expression include immunoassays, such as the enzyme linked immunosorbent assay (ELISA) and the radioimmunoassay (RIA). Suitable antibody assay labels are known in the art and include enzyme labels, such as, glucose oxidase; radioisotopes, such as iodine (^{131}I , ^{125}I , ^{123}I , ^{121}I), carbon (^{14}C), sulfur (^{35}S), tritium (^3H), indium ($^{115\text{m}}\text{In}$, $^{113\text{m}}\text{In}$, ^{112}In , ^{111}In), and technetium (^{99}Tc , $^{99\text{m}}\text{Tc}$), thallium (^{201}Tl), gallium (^{68}Ga , ^{67}Ga), palladium (^{103}Pd), molybdenum (^{99}Mo), xenon (^{133}Xe), fluorine (^{18}F), ^{153}Sm , ^{177}Lu , ^{159}Gd , ^{149}Pm , ^{140}La , ^{175}Yb , ^{166}Ho , ^{90}Y , ^{47}Sc , ^{186}Re , ^{188}Re , ^{142}Pr , ^{105}Rh , ^{97}Ru ; luminescent labels, such as luminol; and fluorescent labels, such as fluorescein and rhodamine, and biotin.

[0358] In addition to assaying levels of polypeptide of the present invention in a biological sample, proteins can also be detected *in vivo* by imaging. Antibody labels or markers for *in vivo* imaging of protein include those detectable by X-radiography, NMR or ESR. For X-radiography, suitable labels include radioisotopes such as barium or cesium, which emit detectable radiation but are not overtly harmful to the subject. Suitable markers for NMR and ESR include those with a detectable characteristic spin, such as deuterium, which may be incorporated into the antibody by labeling of nutrients for the relevant hybridoma.

[0359] A protein-specific antibody or antibody fragment which has been labeled with an appropriate detectable imaging moiety, such as a radioisotope (for example, ^{131}I , ^{112}In , $^{99\text{m}}\text{Tc}$, ^{131}I , ^{125}I , ^{123}I , ^{121}I), carbon (^{14}C), sulfur (^{35}S), tritium (^3H), indium ($^{115\text{m}}\text{In}$, $^{113\text{m}}\text{In}$,

from the sequences of the present invention, specific to tissues, including but not limited to those shown in Table 1. Panels of such reagents can identify tissue by species and/or by organ type. In a similar fashion, these reagents can be used to screen tissue cultures for contamination. Additional non-limiting examples of such uses are further described herein.

[0352] The polynucleotides of the present invention are also useful as hybridization probes for differential identification of the tissue(s) or cell type(s) present in a biological sample. Similarly, polypeptides and antibodies directed to polypeptides of the present invention are useful to provide immunological probes for differential identification of the tissue(s) (e.g., immunohistochemistry assays) or cell type(s) (e.g., immunocytochemistry assays). In addition, for a number of disorders of the above tissues or cells, significantly higher or lower levels of gene expression of the polynucleotides/polypeptides of the present invention may be detected in certain tissues (e.g., tissues expressing polypeptides and/or polynucleotides of the present invention, for example, those disclosed in column 8 of Table 1, and/or cancerous and/or wounded tissues) or bodily fluids (e.g., semen, lymph, vaginal pool, serum, plasma, urine, synovial fluid or spinal fluid) taken from an individual having such a disorder, relative to a "standard" gene expression level, i.e., the expression level in healthy tissue from an individual not having the disorder.

[0353] Thus, the invention provides a diagnostic method of a disorder, which involves: (a) assaying gene expression level in cells or body fluid of an individual; (b) comparing the gene expression level with a standard gene expression level, whereby an increase or decrease in the assayed gene expression level compared to the standard expression level is indicative of a disorder.

[0354] In the very least, the polynucleotides of the present invention can be used as molecular weight markers on Southern gels, as diagnostic probes for the presence of a specific mRNA in a particular cell type, as a probe to "subtract-out" known sequences in the process of discovering novel polynucleotides, for selecting and making oligomers for attachment to a "gene chip" or other support, to raise anti-DNA antibodies using DNA immunization techniques, and as an antigen to elicit an immune response.

herein (see, e.g., the sections labeled "Gene Therapy Methods", and Examples 16, 17 and 18).

[0348] The polynucleotides are also useful for identifying individuals from minute biological samples. The United States military, for example, is considering the use of restriction fragment length polymorphism (RFLP) for identification of its personnel. In this technique, an individual's genomic DNA is digested with one or more restriction enzymes, and probed on a Southern blot to yield unique bands for identifying personnel. This method does not suffer from the current limitations of "Dog Tags" which can be lost, switched, or stolen, making positive identification difficult. The polynucleotides of the present invention can be used as additional DNA markers for RFLP.

[0349] The polynucleotides of the present invention can also be used as an alternative to RFLP, by determining the actual base-by-base DNA sequence of selected portions of an individual's genome. These sequences can be used to prepare PCR primers for amplifying and isolating such selected DNA, which can then be sequenced. Using this technique, individuals can be identified because each individual will have a unique set of DNA sequences. Once an unique ID database is established for an individual, positive identification of that individual, living or dead, can be made from extremely small tissue samples.

[0350] Forensic biology also benefits from using DNA-based identification techniques as disclosed herein. DNA sequences taken from very small biological samples such as tissues, e.g., hair or skin, or body fluids, e.g., blood, saliva, semen, synovial fluid, amniotic fluid, breast milk, lymph, pulmonary sputum or surfactant, urine, fecal matter, etc., can be amplified using PCR. In one prior art technique, gene sequences amplified from polymorphic loci, such as DQa class II HLA gene, are used in forensic biology to identify individuals. (Erlich, H., PCR Technology, Freeman and Co. (1992)). Once these specific polymorphic loci are amplified, they are digested with one or more restriction enzymes, yielding an identifying set of bands on a Southern blot probed with DNA corresponding to the DQa class II HLA gene. Similarly, polynucleotides of the present invention can be used as polymorphic markers for forensic purposes.

[0351] There is also a need for reagents capable of identifying the source of a particular tissue. Such need arises, for example, in forensics when presented with tissue of unknown origin. Appropriate reagents can comprise, for example, DNA probes or primers prepared

hematopoietic origin, in light of the numerous cells and cell types of varying origins which are known to exhibit proliferative phenotypes.

[0346] In addition to the foregoing, a polynucleotide of the present invention can be used to control gene expression through triple helix formation or through antisense DNA or RNA. Antisense techniques are discussed, for example, in Okano, J. *Neurochem.* 56: 560 (1991); "Oligodeoxynucleotides as Antisense Inhibitors of Gene Expression, CRC Press, Boca Raton, FL (1988). Triple helix formation is discussed in, for instance Lee et al., *Nucleic Acids Research* 6: 3073 (1979); Cooney et al., *Science* 241: 456 (1988); and Dervan et al., *Science* 251: 1360 (1991). Both methods rely on binding of the polynucleotide to a complementary DNA or RNA. For these techniques, preferred polynucleotides are usually oligonucleotides 20 to 40 bases in length and complementary to either the region of the gene involved in transcription (triple helix - see Lee et al., *Nucl. Acids Res.* 6:3073 (1979); Cooney et al., *Science* 241:456 (1988); and Dervan et al., *Science* 251:1360 (1991)) or to the mRNA itself (antisense - Okano, J. *Neurochem.* 56:560 (1991); *Oligodeoxy-nucleotides as Antisense Inhibitors of Gene Expression*, CRC Press, Boca Raton, FL (1988)). Triple helix formation optimally results in a shut-off of RNA transcription from DNA, while antisense RNA hybridization blocks translation of an mRNA molecule into polypeptide. The oligonucleotide described above can also be delivered to cells such that the antisense RNA or DNA may be expressed *in vivo* to inhibit production of polypeptide of the present invention antigens. Both techniques are effective in model systems, and the information disclosed herein can be used to design antisense or triple helix polynucleotides in an effort to treat disease, and in particular, for the treatment of proliferative diseases and/or conditions. Non-limiting antisense and triple helix methods encompassed by the present invention are more thoroughly described elsewhere herein (see, e.g., the section labeled "Antisense and Ribozyme (Antagonists)").

[0347] Polynucleotides of the present invention are also useful in gene therapy. One goal of gene therapy is to insert a normal gene into an organism having a defective gene, in an effort to correct the genetic defect. The polynucleotides disclosed in the present invention offer a means of targeting such genetic defects in a highly accurate manner. Another goal is to insert a new gene that was not present in the host genome, thereby producing a new trait in the host cell. Additional non-limiting examples of gene therapy methods encompassed by the present invention are more thoroughly described elsewhere

[0343] The compounds of the present invention have uses which include, but are not limited to, detecting cancer in mammals. In particular the invention is useful during diagnosis of pathological cell proliferative neoplasias which include, but are not limited to: acute myelogenous leukemias including acute monocytic leukemia, acute myeloblastic leukemia, acute promyelocytic leukemia, acute myelomonocytic leukemia, acute erythroleukemia, acute megakaryocytic leukemia, and acute undifferentiated leukemia, etc.; and chronic myelogenous leukemias including chronic myelomonocytic leukemia, chronic granulocytic leukemia, etc. Preferred mammals include monkeys, apes, cats, dogs, cows, pigs, horses, rabbits and humans. Particularly preferred are humans.

[0344] Pathological cell proliferative disorders are often associated with inappropriate activation of proto-oncogenes. (Gelmann, E. P. et al., "The Etiology of Acute Leukemia: Molecular Genetics and Viral Oncology," in Neoplastic Diseases of the Blood, Vol 1., Wiernik, P. H. et al. eds., 161-182 (1985)). Neoplasias are now believed to result from the qualitative alteration of a normal cellular gene product, or from the quantitative modification of gene expression by insertion into the chromosome of a viral sequence, by chromosomal translocation of a gene to a more actively transcribed region, or by some other mechanism. (Gelmann et al., *supra*) It is likely that mutated or altered expression of specific genes is involved in the pathogenesis of some leukemias, among other tissues and cell types. (Gelmann et al., *supra*) Indeed, the human counterparts of the oncogenes involved in some animal neoplasias have been amplified or translocated in some cases of human leukemia and carcinoma. (Gelmann et al., *supra*)

[0345] For example, c-myc expression is highly amplified in the non-lymphocytic leukemia cell line HL-60. When HL-60 cells are chemically induced to stop proliferation, the level of c-myc is found to be downregulated. (International Publication Number WO 91/15580). However, it has been shown that exposure of HL-60 cells to a DNA construct that is complementary to the 5' end of c-myc or c-myb blocks translation of the corresponding mRNAs which downregulates expression of the c-myc or c-myb proteins and causes arrest of cell proliferation and differentiation of the treated cells. (International Publication Number WO 91/15580; Wickstrom et al., Proc. Natl. Acad. Sci. 85:1028 (1988); Anfossi et al., Proc. Natl. Acad. Sci. 86:3379 (1989)). However, the skilled artisan would appreciate the present invention's usefulness is not be limited to treatment, prevention, and/or prognosis of proliferative disorders of cells and tissues of

or a "biological chip" as described in US Patents 5,837,832, 5,874,219, and 5,856,174. Further, such a gene chip with polynucleotides of the invention attached may be used to identify polymorphisms between the isolated polynucleotide sequences of the invention, with polynucleotides isolated from a test subject. The knowledge of such polymorphisms (i.e. their location, as well as, their existence) would be beneficial in identifying disease loci for many disorders, such as for example, in neural disorders, immune system disorders, muscular disorders, reproductive disorders, gastrointestinal disorders, pulmonary disorders, digestive disorders, metabolic disorders, cardiovascular disorders, renal disorders, proliferative disorders, and/or cancerous diseases and conditions. Such a method is described in US Patents 5,858,659 and 5,856,104. The US Patents referenced *supra* are hereby incorporated by reference in their entirety herein.

[0342] The present invention encompasses polynucleotides of the present invention that are chemically synthesized, or reproduced as peptide nucleic acids (PNA), or according to other methods known in the art. The use of PNAs would serve as the preferred form if the polynucleotides of the invention are incorporated onto a solid support, or gene chip. For the purposes of the present invention, a peptide nucleic acid (PNA) is a polyamide type of DNA analog and the monomeric units for adenine, guanine, thymine and cytosine are available commercially (Perceptive Biosystems). Certain components of DNA, such as phosphorus, phosphorus oxides, or deoxyribose derivatives, are not present in PNAs. As disclosed by Nielsen et al., Science 254, 1497 (1991); and Egholm et al., Nature 365, 666 (1993), PNAs bind specifically and tightly to complementary DNA strands and are not degraded by nucleases. In fact, PNA binds more strongly to DNA than DNA itself does. This is probably because there is no electrostatic repulsion between the two strands, and also the polyamide backbone is more flexible. Because of this, PNA/DNA duplexes bind under a wider range of stringency conditions than DNA/DNA duplexes, making it easier to perform multiplex hybridization. Smaller probes can be used than with DNA due to the strong binding. In addition, it is more likely that single base mismatches can be determined with PNA/DNA hybridization because a single mismatch in a PNA/DNA 15-mer lowers the melting point ($T_{sub.m}$) by 8°-20° C, vs. 4°-16° C for the DNA/DNA 15-mer duplex. Also, the absence of charge groups in PNA means that hybridization can be done at low ionic strengths and reduce possible interference by salt during the analysis.

polynucleotide probes defining an internal region of the polynucleotide of the invention, where each probe has one strand containing a 31'-mer-end internal to the region. In a further embodiment, the probes may be useful as primers for polymerase chain reaction amplification.

[0338] Where a diagnosis of a related disorder, including, for example, diagnosis of a tumor, has already been made according to conventional methods, the present invention is useful as a prognostic indicator, whereby patients exhibiting enhanced or depressed polynucleotide of the invention expression will experience a worse clinical outcome relative to patients expressing the gene at a level nearer the standard level.

[0339] By "measuring the expression level of polynucleotides of the invention" is intended qualitatively or quantitatively measuring or estimating the level of the polypeptide of the invention or the level of the mRNA encoding the polypeptide of the invention in a first biological sample either directly (e.g., by determining or estimating absolute protein level or mRNA level) or relatively (e.g., by comparing to the polypeptide level or mRNA level in a second biological sample). Preferably, the polypeptide level or mRNA level in the first biological sample is measured or estimated and compared to a standard polypeptide level or mRNA level, the standard being taken from a second biological sample obtained from an individual not having the related disorder or being determined by averaging levels from a population of individuals not having a related disorder. As will be appreciated in the art, once a standard polypeptide level or mRNA level is known, it can be used repeatedly as a standard for comparison.

[0340] By "biological sample" is intended any biological sample obtained from an individual, body fluid, cell line, tissue culture, or other source which contains polypeptide of the present invention or the corresponding mRNA. As indicated, biological samples include body fluids (such as semen, lymph, vaginal pool, sera, plasma, urine, synovial fluid and spinal fluid) which contain the polypeptide of the present invention, and tissue sources found to express the polypeptide of the present invention. Methods for obtaining tissue biopsies and body fluids from mammals are well known in the art. Where the biological sample is to include mRNA, a tissue biopsy is the preferred source.

[0341] The method(s) provided above may preferably be applied in a diagnostic method and/or kits in which polynucleotides and/or polypeptides of the invention are attached to a solid support. In one exemplary method, the support may be a "gene chip"

chromosomal region associated with the disease could be one of 50-500 potential causative genes.

[0334] Thus, once coinheritance is established, differences in a polynucleotide of the invention and the corresponding gene between affected and unaffected individuals can be examined. First, visible structural alterations in the chromosomes, such as deletions or translocations, are examined in chromosome spreads or by PCR. If no structural alterations exist, the presence of point mutations are ascertained. Mutations observed in some or all affected individuals, but not in normal individuals, indicates that the mutation may cause the disease. However, complete sequencing of the polypeptide and the corresponding gene from several normal individuals is required to distinguish the mutation from a polymorphism. If a new polymorphism is identified, this polymorphic polypeptide can be used for further linkage analysis.

[0335] Furthermore, increased or decreased expression of the gene in affected individuals as compared to unaffected individuals can be assessed using the polynucleotides of the invention. Any of these alterations (altered expression, chromosomal rearrangement, or mutation) can be used as a diagnostic or prognostic marker. Diagnostic and prognostic methods, kits and reagents encompassed by the present invention are briefly described below and more thoroughly elsewhere herein (see e.g., the sections labeled "Antibodies", "Diagnostic Assays", and "Methods for Detecting Diseases").

[0336] Thus, the invention also provides a diagnostic method useful during diagnosis of a disorder, involving measuring the expression level of polynucleotides of the present invention in cells or body fluid from an individual and comparing the measured gene expression level with a standard level of polynucleotide expression level, whereby an increase or decrease in the gene expression level compared to the standard is indicative of a disorder. Additional non-limiting examples of diagnostic methods encompassed by the present invention are more thoroughly described elsewhere herein (see, e.g., Example 12).

[0337] In still another embodiment, the invention includes a kit for analyzing samples for the presence of proliferative and/or cancerous polynucleotides derived from a test subject. In a general embodiment, the kit includes at least one polynucleotide probe containing a nucleotide sequence that will specifically hybridize with a polynucleotide of the invention and a suitable container. In a specific embodiment, the kit includes two

chromosomes, preselection by hybridization to construct chromosome specific-cDNA libraries, and computer mapping techniques (See, e.g., Shuler, Trends Biotechnol 16:456-459 (1998) which is hereby incorporated by reference in its entirety).

[0329] Precise chromosomal location of the polynucleotides can also be achieved using fluorescence in situ hybridization (FISH) of a metaphase chromosomal spread. This technique uses polynucleotides as short as 500 or 600 bases; however, polynucleotides 2,000-4,000 bp are preferred. For a review of this technique, see Verma et al., "Human Chromosomes: a Manual of Basic Techniques," Pergamon Press, New York (1988).

[0330] For chromosome mapping, the polynucleotides can be used individually (to mark a single chromosome or a single site on that chromosome) or in panels (for marking multiple sites and/or multiple chromosomes).

[0331] Thus, the present invention also provides a method for chromosomal localization which involves (a) preparing PCR primers from the polynucleotide sequences in Table 1 and/or Table 2 and SEQ ID NO:X and (b) screening somatic cell hybrids containing individual chromosomes.

[0332] The polynucleotides of the present invention would likewise be useful for radiation hybrid mapping, HAPPY mapping, and long range restriction mapping. For a review of these techniques and others known in the art, see, e.g. Dear, "Genome Mapping: A Practical Approach," IRL Press at Oxford University Press, London (1997); Aydin, J. Mol. Med. 77:691-694 (1999); Hacia et al., Mol. Psychiatry 3:483-492 (1998); Herrick et al., Chromosome Res. 7:409-423 (1999); Hamilton et al., Methods Cell Biol. 62:265-280 (2000); and/or Ott, J. Hered. 90:68-70 (1999) each of which is hereby incorporated by reference in its entirety.

[0333] Once a polynucleotide has been mapped to a precise chromosomal location, the physical position of the polynucleotide can be used in linkage analysis. Linkage analysis establishes coinheritance between a chromosomal location and presentation of a particular disease. (Disease mapping data are found, for example, in V. McKusick, Mendelian Inheritance in Man (available on line through Johns Hopkins University Welch Medical Library)). Column 10 of Table 1 provides an OMIM reference identification number of diseases associated with the cytologic band disclosed in column 9 of Table 1, as determined using techniques described herein and by reference to Table 5. Assuming 1 megabase mapping resolution and one gene per 20 kb, a cDNA precisely localized to a

such as an activated carboxyl, hydroxyl, or aldehyde group. Alternatively, streptavidin coated plates can be used in conjunction with biotinylated antigen(s).

[0324] Thus, the invention provides an assay system or kit for carrying out this diagnostic method. The kit generally includes a support with surface-bound recombinant antigens, and a reporter-labeled anti-human antibody for detecting surface-bound anti-antigen antibody.

Uses of the Polynucleotides

[0325] Each of the polynucleotides identified herein can be used in numerous ways as reagents. The following description should be considered exemplary and utilizes known techniques.

[0326] The polynucleotides of the present invention are useful for chromosome identification. There exists an ongoing need to identify new chromosome markers, since few chromosome marking reagents, based on actual sequence data (repeat polymorphisms), are presently available. Each sequence is specifically targeted to and can hybridize with a particular location on an individual human chromosome, thus each polynucleotide of the present invention can routinely be used as a chromosome marker using techniques known in the art. Table 1, column 9 provides the chromosome location of some of the polynucleotides of the invention.

[0327] Briefly, sequences can be mapped to chromosomes by preparing PCR primers (preferably at least 15 bp (e.g., 15-25 bp) from the sequences shown in SEQ ID NO:X. Primers can optionally be selected using computer analysis so that primers do not span more than one predicted exon in the genomic DNA. These primers are then used for PCR screening of somatic cell hybrids containing individual human chromosomes. Only those hybrids containing the human gene corresponding to SEQ ID NO:X will yield an amplified fragment.

[0328] Similarly, somatic hybrids provide a rapid method of PCR mapping the polynucleotides to particular chromosomes. Three or more clones can be assigned per day using a single thermal cycler. Moreover, sublocalization of the polynucleotides can be achieved with panels of specific chromosome fragments. Other gene mapping strategies that can be used include in situ hybridization, prescreening with labeled flow-sorted

to a fluorescent compound such as fluorescein or rhodamine which can be detected by flow cytometry). In specific embodiments, the kit may include a recombinantly produced or chemically synthesized polypeptide antigen. The polypeptide antigen of the kit may also be attached to a solid support.

[0320] In a more specific embodiment the detecting means of the above-described kit includes a solid support to which said polypeptide antigen is attached. Such a kit may also include a non-attached reporter-labeled anti-human antibody. In this embodiment, binding of the antibody to the polypeptide antigen can be detected by binding of the said reporter-labeled antibody.

[0321] In an additional embodiment, the invention includes a diagnostic kit for use in screening serum containing antigens of the polypeptide of the invention. The diagnostic kit includes a substantially isolated antibody specifically immunoreactive with polypeptide or polynucleotide antigens, and means for detecting the binding of the polynucleotide or polypeptide antigen to the antibody. In one embodiment, the antibody is attached to a solid support. In a specific embodiment, the antibody may be a monoclonal antibody. The detecting means of the kit may include a second, labeled monoclonal antibody. Alternatively, or in addition, the detecting means may include a labeled, competing antigen.

[0322] In one diagnostic configuration, test serum is reacted with a solid phase reagent having a surface-bound antigen obtained by the methods of the present invention. After binding with specific antigen antibody to the reagent and removing unbound serum components by washing, the reagent is reacted with reporter-labeled anti-human antibody to bind reporter to the reagent in proportion to the amount of bound anti-antigen antibody on the solid support. The reagent is again washed to remove unbound labeled antibody, and the amount of reporter associated with the reagent is determined. Typically, the reporter is an enzyme which is detected by incubating the solid phase in the presence of a suitable fluorometric, luminescent or colorimetric substrate (Sigma, St. Louis, MO).

[0323] The solid surface reagent in the above assay is prepared by known techniques for attaching protein material to solid support material, such as polymeric beads, dip sticks, 96-well plate or filter material. These attachment methods generally include non-specific adsorption of the protein to the support or covalent attachment of the protein, typically through a free amine group, to a chemically reactive group on the solid support,

label. Methods and devices that may be used in the diagnostic methods of the invention include, but are not limited to, computed tomography (CT), whole body scan such as position emission tomography (PET), magnetic resonance imaging (MRI), and sonography.

[0317] In a specific embodiment, the molecule is labeled with a radioisotope and is detected in the patient using a radiation responsive surgical instrument (Thurston et al., U.S. Patent No. 5,441,050). In another embodiment, the molecule is labeled with a fluorescent compound and is detected in the patient using a fluorescence responsive scanning instrument. In another embodiment, the molecule is labeled with a positron emitting metal and is detected in the patient using positron emission-tomography. In yet another embodiment, the molecule is labeled with a paramagnetic label and is detected in a patient using magnetic resonance imaging (MRI).

Kits

[0318] The present invention provides kits that can be used in the above methods. In one embodiment, a kit comprises an antibody of the invention, preferably a purified antibody, in one or more containers. In a specific embodiment, the kits of the present invention contain a substantially isolated polypeptide comprising an epitope which is specifically immunoreactive with an antibody included in the kit. Preferably, the kits of the present invention further comprise a control antibody which does not react with the polypeptide of interest. In another specific embodiment, the kits of the present invention contain a means for detecting the binding of an antibody to a polypeptide of interest (e.g., the antibody may be conjugated to a detectable substrate such as a fluorescent compound, an enzymatic substrate, a radioactive compound or a luminescent compound, or a second antibody which recognizes the first antibody may be conjugated to a detectable substrate).

[0319] In another specific embodiment of the present invention, the kit is a diagnostic kit for use in screening serum containing antibodies specific against proliferative and/or cancerous polynucleotides and polypeptides. Such a kit may include a control antibody that does not react with the polypeptide of interest. Such a kit may include a substantially isolated polypeptide antigen comprising an epitope which is specifically immunoreactive with at least one anti-polypeptide antigen antibody. Further, such a kit includes means for detecting the binding of said antibody to the antigen (e.g., the antibody may be conjugated

administering (for example, parenterally, subcutaneously, or intraperitoneally) to a subject an effective amount of a labeled molecule which specifically binds to the polypeptide of interest; b) waiting for a time interval following the administering for permitting the labeled molecule to preferentially concentrate at sites in the subject where the polypeptide is expressed (and for unbound labeled molecule to be cleared to background level); c) determining background level; and d) detecting the labeled molecule in the subject, such that detection of labeled molecule above the background level indicates that the subject has a particular disease or disorder associated with aberrant expression of the polypeptide of interest. Background level can be determined by various methods including, comparing the amount of labeled molecule detected to a standard value previously determined for a particular system.

[0313] It will be understood in the art that the size of the subject and the imaging system used will determine the quantity of imaging moiety needed to produce diagnostic images. In the case of a radioisotope moiety, for a human subject, the quantity of radioactivity injected will normally range from about 5 to 20 millicuries of ^{99m}Tc. The labeled antibody or antibody fragment will then preferentially accumulate at the location of cells which contain the specific protein. *In vivo* tumor imaging is described in S.W. Burchiel et al., "Immunopharmacokinetics of Radiolabeled Antibodies and Their Fragments." (Chapter 13 in *Tumor Imaging: The Radiochemical Detection of Cancer*, S.W. Burchiel and B. A. Rhodes, eds., Masson Publishing Inc. (1982)).

[0314] Depending on several variables, including the type of label used and the mode of administration, the time interval following the administration for permitting the labeled molecule to preferentially concentrate at sites in the subject and for unbound labeled molecule to be cleared to background level is 6 to 48 hours or 6 to 24 hours or 6 to 12 hours. In another embodiment the time interval following administration is 5 to 20 days or 5 to 10 days.

[0315] In an embodiment, monitoring of the disease or disorder is carried out by repeating the method for diagnosing the disease or disorder, for example, one month after initial diagnosis, six months after initial diagnosis, one year after initial diagnosis, etc.

[0316] Presence of the labeled molecule can be detected in the patient using methods known in the art for *in vivo* scanning. These methods depend upon the type of label used. Skilled artisans will be able to determine the appropriate method for detecting a particular

and/or activity of a polypeptide of the invention. The invention provides for the detection of aberrant expression of a polypeptide of interest, comprising (a) assaying the expression of the polypeptide of interest in cells or body fluid of an individual using one or more antibodies specific to the polypeptide interest and (b) comparing the level of gene expression with a standard gene expression level, whereby an increase or decrease in the assayed polypeptide gene expression level compared to the standard expression level is indicative of aberrant expression.

[0310] The invention provides a diagnostic assay for diagnosing a disorder, comprising (a) assaying the expression of the polypeptide of interest in cells or body fluid of an individual using one or more antibodies specific to the polypeptide interest and (b) comparing the level of gene expression with a standard gene expression level, whereby an increase or decrease in the assayed polypeptide gene expression level compared to the standard expression level is indicative of a particular disorder. With respect to cancer, the presence of a relatively high amount of transcript in biopsied tissue from an individual may indicate a predisposition for the development of the disease, or may provide a means for detecting the disease prior to the appearance of actual clinical symptoms. A more definitive diagnosis of this type may allow health professionals to employ preventative measures or aggressive treatment earlier thereby preventing the development or further progression of the cancer.

[0311] Antibodies of the invention can be used to assay protein levels in a biological sample using classical immunohistological methods known to those of skill in the art (e.g., see Jalkanen et al., J. Cell. Biol. 101:976-985 (1985); Jalkanen et al., J. Cell. Biol. 105:3087-3096 (1987)). Other antibody-based methods useful for detecting protein gene expression include immunoassays, such as the enzyme linked immunosorbent assay (ELISA) and the radioimmunoassay (RIA). Suitable antibody assay labels are known in the art and include enzyme labels, such as, glucose oxidase; radioisotopes, such as iodine (^{125}I , ^{121}I), carbon (^{14}C), sulfur (^{35}S), tritium (^3H), indium (^{112}In), and technetium (^{99}Tc); luminescent labels, such as luminol; and fluorescent labels, such as fluorescein and rhodamine, and biotin.

[0312] One facet of the invention is the detection and diagnosis of a disease or disorder associated with aberrant expression of a polypeptide of interest in an animal, preferably a mammal and most preferably a human. In one embodiment, diagnosis comprises: a)

cations such as those derived from sodium, potassium, ammonium, calcium, ferric hydroxides, isopropylamine, triethylamine, 2-ethylamino ethanol, histidine, procaine, etc.

[0306] The amount of the compound of the invention which will be effective in the treatment, inhibition and prevention of a disease or disorder associated with aberrant expression and/or activity of a polypeptide of the invention can be determined by standard clinical techniques. In addition, in vitro assays may optionally be employed to help identify optimal dosage ranges. The precise dose to be employed in the formulation will also depend on the route of administration, and the seriousness of the disease or disorder, and should be decided according to the judgment of the practitioner and each patient's circumstances. Effective doses may be extrapolated from dose-response curves derived from in vitro or animal model test systems.

[0307] For antibodies, the dosage administered to a patient is typically 0.1 mg/kg to 100 mg/kg of the patient's body weight. Preferably, the dosage administered to a patient is between 0.1 mg/kg and 20 mg/kg of the patient's body weight, more preferably 1 mg/kg to 10 mg/kg of the patient's body weight. Generally, human antibodies have a longer half-life within the human body than antibodies from other species due to the immune response to the foreign polypeptides. Thus, lower dosages of human antibodies and less frequent administration is often possible. Further, the dosage and frequency of administration of antibodies of the invention may be reduced by enhancing uptake and tissue penetration (e.g., into the brain) of the antibodies by modifications such as, for example, lipidation.

[0308] The invention also provides a pharmaceutical pack or kit comprising one or more containers filled with one or more of the ingredients of the pharmaceutical compositions of the invention. Optionally associated with such container(s) can be a notice in the form prescribed by a governmental agency regulating the manufacture, use or sale of pharmaceuticals or biological products, which notice reflects approval by the agency of manufacture, use or sale for human administration.

Diagnosis and Imaging

[0309] Labeled antibodies, and derivatives and analogs thereof, which specifically bind to a polypeptide of interest can be used for diagnostic purposes to detect, diagnose, or monitor diseases, disorders, and/or conditions associated with the aberrant expression

injectable solutions. Suitable pharmaceutical excipients include starch, glucose, lactose, sucrose, gelatin, malt, rice, flour, chalk, silica gel, sodium stearate, glycerol monostearate, talc, sodium chloride, dried skim milk, glycerol, propylene, glycol, water, ethanol and the like. The composition, if desired, can also contain minor amounts of wetting or emulsifying agents, or pH buffering agents. These compositions can take the form of solutions, suspensions, emulsion, tablets, pills, capsules, powders, sustained-release formulations and the like. The composition can be formulated as a suppository, with traditional binders and carriers such as triglycerides. Oral formulation can include standard carriers such as pharmaceutical grades of mannitol, lactose, starch, magnesium stearate, sodium saccharine, cellulose, magnesium carbonate, etc. Examples of suitable pharmaceutical carriers are described in "Remington's Pharmaceutical Sciences" by E.W. Martin. Such compositions will contain a therapeutically effective amount of the compound, preferably in purified form, together with a suitable amount of carrier so as to provide the form for proper administration to the patient. The formulation should suit the mode of administration.

[0304] In a preferred embodiment, the composition is formulated in accordance with routine procedures as a pharmaceutical composition adapted for intravenous administration to human beings. Typically, compositions for intravenous administration are solutions in sterile isotonic aqueous buffer. Where necessary, the composition may also include a solubilizing agent and a local anesthetic such as lignocaine to ease pain at the site of the injection. Generally, the ingredients are supplied either separately or mixed together in unit dosage form, for example, as a dry lyophilized powder or water free concentrate in a hermetically sealed container such as an ampoule or sachette indicating the quantity of active agent. Where the composition is to be administered by infusion, it can be dispensed with an infusion bottle containing sterile pharmaceutical grade water or saline. Where the composition is administered by injection, an ampoule of sterile water for injection or saline can be provided so that the ingredients may be mixed prior to administration.

[0305] The compounds of the invention can be formulated as neutral or salt forms. Pharmaceutically acceptable salts include those formed with anions such as those derived from hydrochloric, phosphoric, acetic, oxalic, tartaric acids, etc., and those formed with

Drug Product Design and Performance, Smolen and Ball (eds.), Wiley, New York (1984); Ranger and Peppas, J., *Macromol. Sci. Rev. Macromol. Chem.* 23:61 (1983); see also Levy et al., *Science* 228:190 (1985); During et al., *Ann. Neurol.* 25:351 (1989); Howard et al., *J. Neurosurg.* 71:105 (1989)). In yet another embodiment, a controlled release system can be placed in proximity of the therapeutic target, e.g., the brain, thus requiring only a fraction of the systemic dose (see, e.g., Goodson, in *Medical Applications of Controlled Release*, *supra*, vol. 2, pp. 115-138 (1984)).

[0301] Other controlled release systems are discussed in the review by Langer (*Science* 249:1527-1533 (1990)).

[0302] In a specific embodiment where the compound of the invention is a nucleic acid encoding a protein, the nucleic acid can be administered *in vivo* to promote expression of its encoded protein, by constructing it as part of an appropriate nucleic acid expression vector and administering it so that it becomes intracellular, e.g., by use of a retroviral vector (see U.S. Patent No. 4,980,286), or by direct injection, or by use of microparticle bombardment (e.g., a gene gun; Biolistic, Dupont), or coating with lipids or cell-surface receptors or transfecting agents, or by administering it in linkage to a homeobox-like peptide which is known to enter the nucleus (see e.g., Joliot et al., *Proc. Natl. Acad. Sci. USA* 88:1864-1868 (1991)), etc. Alternatively, a nucleic acid can be introduced intracellularly and incorporated within host cell DNA for expression, by homologous recombination.

[0303] The present invention also provides pharmaceutical compositions. Such compositions comprise a therapeutically effective amount of a compound, and a pharmaceutically acceptable carrier. In a specific embodiment, the term "pharmaceutically acceptable" means approved by a regulatory agency of the Federal or a state government or listed in the U.S. Pharmacopeia or other generally recognized pharmacopeia for use in animals, and more particularly in humans. The term "carrier" refers to a diluent, adjuvant, excipient, or vehicle with which the therapeutic is administered. Such pharmaceutical carriers can be sterile liquids, such as water and oils, including those of petroleum, animal, vegetable or synthetic origin, such as peanut oil, soybean oil, mineral oil, sesame oil and the like. Water is a preferred carrier when the pharmaceutical composition is administered intravenously. Saline solutions and aqueous dextrose and glycerol solutions can also be employed as liquid carriers, particularly for

limited to intradermal, intramuscular, intraperitoneal, intravenous, subcutaneous, intranasal, epidural, and oral routes. The compounds or compositions may be administered by any convenient route, for example by infusion or bolus injection, by absorption through epithelial or mucocutaneous linings (e.g., oral mucosa, rectal and intestinal mucosa, etc.) and may be administered together with other biologically active agents. Administration can be systemic or local. In addition, it may be desirable to introduce the pharmaceutical compounds or compositions of the invention into the central nervous system by any suitable route, including intraventricular and intrathecal injection; intraventricular injection may be facilitated by an intraventricular catheter, for example, attached to a reservoir, such as an Ommaya reservoir. Pulmonary administration can also be employed, e.g., by use of an inhaler or nebulizer, and formulation with an aerosolizing agent.

[0298] In a specific embodiment, it may be desirable to administer the pharmaceutical compounds or compositions of the invention locally to the area in need of treatment; this may be achieved by, for example, and not by way of limitation, local infusion during surgery, topical application, e.g., in conjunction with a wound dressing after surgery, by injection, by means of a catheter, by means of a suppository, or by means of an implant, said implant being of a porous, non-porous, or gelatinous material, including membranes, such as sialastic membranes, or fibers. Preferably, when administering a protein, including an antibody, of the invention, care must be taken to use materials to which the protein does not absorb.

[0299] In another embodiment, the compound or composition can be delivered in a vesicle, in particular a liposome (see Langer, *Science* 249:1527-1533 (1990); Treat et al., in *Liposomes in the Therapy of Infectious Disease and Cancer*, Lopez-Berestein and Fidler (eds.), Liss, New York, pp. 353- 365 (1989); Lopez-Berestein, *ibid.*, pp. 317-327; see generally *ibid.*)

[0300] In yet another embodiment, the compound or composition can be delivered in a controlled release system. In one embodiment, a pump may be used (see Langer, *supra*; Sefton, *CRC Crit. Ref. Biomed. Eng.* 14:201 (1987); Buchwald et al., *Surgery* 88:507 (1980); Saudek et al., *N. Engl. J. Med.* 321:574 (1989)). In another embodiment, polymeric materials can be used (see *Medical Applications of Controlled Release*, Langer and Wise (eds.), CRC Pres., Boca Raton, Florida (1974); *Controlled Drug Bioavailability*,

Demonstration of Therapeutic or Prophylactic Activity

[0293] The compounds or pharmaceutical compositions of the invention are preferably tested *in vitro*, and then *in vivo* for the desired therapeutic or prophylactic activity, prior to use in humans. For example, *in vitro* assays to demonstrate the therapeutic or prophylactic utility of a compound or pharmaceutical composition include, the effect of a compound on a cell line or a patient tissue sample. The effect of the compound or composition on the cell line and/or tissue sample can be determined utilizing techniques known to those of skill in the art including, but not limited to, rosette formation assays and cell lysis assays. In accordance with the invention, *in vitro* assays which can be used to determine whether administration of a specific compound is indicated, include *in vitro* cell culture assays in which a patient tissue sample is grown in culture, and exposed to or otherwise administered a compound, and the effect of such compound upon the tissue sample is observed.

[0294] Therapeutic/Prophylactic Administration and Composition

[0295] The invention provides methods of treatment, inhibition and prophylaxis by administration to a subject of an effective amount of a compound or pharmaceutical composition of the invention, preferably a polypeptide or antibody of the invention. In a preferred embodiment, the compound is substantially purified (e.g., substantially free from substances that limit its effect or produce undesired side-effects). The subject is preferably an animal, including but not limited to animals such as cows, pigs, horses, chickens, cats, dogs, etc., and is preferably a mammal, and most preferably human.

[0296] Formulations and methods of administration that can be employed when the compound comprises a nucleic acid or an immunoglobulin are described above; additional appropriate formulations and routes of administration can be selected from among those described herein below.

[0297] Various delivery systems are known and can be used to administer a compound of the invention, e.g., encapsulation in liposomes, microparticles, microcapsules, recombinant cells capable of expressing the compound, receptor-mediated endocytosis (see, e.g., Wu and Wu, J. Biol. Chem. 262:4429-4432 (1987)), construction of a nucleic acid as part of a retroviral or other vector, etc. Methods of introduction include but are not

Pharmac. Ther. 29:69-92m (1985) and may be used in accordance with the present invention, provided that the necessary developmental and physiological functions of the recipient cells are not disrupted. The technique should provide for the stable transfer of the nucleic acid to the cell, so that the nucleic acid is expressible by the cell and preferably heritable and expressible by its cell progeny.

[0288] The resulting recombinant cells can be delivered to a patient by various methods known in the art. Recombinant blood cells (e.g., hematopoietic stem or progenitor cells) are preferably administered intravenously. The amount of cells envisioned for use depends on the desired effect, patient state, etc., and can be determined by one skilled in the art.

[0289] Cells into which a nucleic acid can be introduced for purposes of gene therapy encompass any desired, available cell type, and include but are not limited to epithelial cells, endothelial cells, keratinocytes, fibroblasts, muscle cells, hepatocytes; blood cells such as T lymphocytes, B lymphocytes, monocytes, macrophages, neutrophils, eosinophils, megakaryocytes, granulocytes; various stem or progenitor cells, in particular hematopoietic stem or progenitor cells, e.g., as obtained from bone marrow, umbilical cord blood, peripheral blood, fetal liver, etc.

[0290] In a preferred embodiment, the cell used for gene therapy is autologous to the patient.

[0291] In an embodiment in which recombinant cells are used in gene therapy, nucleic acid sequences encoding an antibody are introduced into the cells such that they are expressible by the cells or their progeny, and the recombinant cells are then administered *in vivo* for therapeutic effect. In a specific embodiment, stem or progenitor cells are used. Any stem and/or progenitor cells which can be isolated and maintained *in vitro* can potentially be used in accordance with this embodiment of the present invention (see e.g. PCT Publication WO 94/08598; Stemple and Anderson, Cell 71:973-985 (1992); Rheinwald, Meth. Cell Bio. 21A:229 (1980); and Pittelkow and Scott, Mayo Clinic Proc. 61:771 (1986)).

[0292] In a specific embodiment, the nucleic acid to be introduced for purposes of gene therapy comprises an inducible promoter operably linked to the coding region, such that expression of the nucleic acid is controllable by the presence or absence of an appropriate inducer of transcription.

(1994); Salmons and Gunzberg, Human Gene Therapy 4:129-141 (1993); and Grossman and Wilson, Curr. Opin. in Genetics and Devel. 3:110-114 (1993).

[0284] Adenoviruses are other viral vectors that can be used in gene therapy. Adenoviruses are especially attractive vehicles for delivering genes to respiratory epithelia. Adenoviruses naturally infect respiratory epithelia where they cause a mild disease. Other targets for adenovirus-based delivery systems are liver, the central nervous system, endothelial cells, and muscle. Adenoviruses have the advantage of being capable of infecting non-dividing cells. Kozarsky and Wilson, Current Opinion in Genetics and Development 3:499-503 (1993) present a review of adenovirus-based gene therapy. Bout et al., Human Gene Therapy 5:3-10 (1994) demonstrated the use of adenovirus vectors to transfer genes to the respiratory epithelia of rhesus monkeys. Other instances of the use of adenoviruses in gene therapy can be found in Rosenfeld et al., Science 252:431-434 (1991); Rosenfeld et al., Cell 68:143-155 (1992); Mastrangeli et al., J. Clin. Invest. 91:225-234 (1993); PCT Publication WO94/12649; and Wang, et al., Gene Therapy 2:775-783 (1995). In a preferred embodiment, adenovirus vectors are used.

[0285] Adeno-associated virus (AAV) has also been proposed for use in gene therapy (Walsh et al., Proc. Soc. Exp. Biol. Med. 204:289-300 (1993); U.S. Patent No. 5,436,146).

[0286] Another approach to gene therapy involves transferring a gene to cells in tissue culture by such methods as electroporation, lipofection, calcium phosphate mediated transfection, or viral infection. Usually, the method of transfer includes the transfer of a selectable marker to the cells. The cells are then placed under selection to isolate those cells that have taken up and are expressing the transferred gene. Those cells are then delivered to a patient.

[0287] In this embodiment, the nucleic acid is introduced into a cell prior to administration *in vivo* of the resulting recombinant cell. Such introduction can be carried out by any method known in the art, including but not limited to transfection, electroporation, microinjection, infection with a viral or bacteriophage vector containing the nucleic acid sequences, cell fusion, chromosome-mediated gene transfer, microcell-mediated gene transfer, spheroplast fusion, etc. Numerous techniques are known in the art for the introduction of foreign genes into cells (see, e.g., Loeffler and Behr, Meth. Enzymol. 217:599-618 (1993); Cohen et al., Meth. Enzymol. 217:618-644 (1993); Cline,

[0282] In a specific embodiment, the nucleic acid sequences are directly administered *in vivo*, where it is expressed to produce the encoded product. This can be accomplished by any of numerous methods known in the art, e.g., by constructing them as part of an appropriate nucleic acid expression vector and administering it so that they become intracellular, e.g., by infection using defective or attenuated retrovirals or other viral vectors (see U.S. Patent No. 4,980,286), or by direct injection of naked DNA, or by use of microparticle bombardment (e.g., a gene gun; Biolistic, Dupont), or coating with lipids or cell-surface receptors or transfecting agents, encapsulation in liposomes, microparticles, or microcapsules, or by administering them in linkage to a peptide which is known to enter the nucleus, by administering it in linkage to a ligand subject to receptor-mediated endocytosis (see, e.g., Wu and Wu, J. Biol. Chem. 262:4429-4432 (1987)) (which can be used to target cell types specifically expressing the receptors), etc. In another embodiment, nucleic acid-ligand complexes can be formed in which the ligand comprises a fusogenic viral peptide to disrupt endosomes, allowing the nucleic acid to avoid lysosomal degradation. In yet another embodiment, the nucleic acid can be targeted *in vivo* for cell specific uptake and expression, by targeting a specific receptor (see, e.g., PCT Publications WO 92/06180; WO 92/22635; WO92/20316; WO93/14188, WO 93/20221). Alternatively, the nucleic acid can be introduced intracellularly and incorporated within host cell DNA for expression, by homologous recombination (Koller and Smithies, Proc. Natl. Acad. Sci. USA 86:8932-8935 (1989); Zijlstra et al., Nature 342:435-438 (1989)).

[0283] In a specific embodiment, viral vectors that contains nucleic acid sequences encoding an antibody of the invention are used. For example, a retroviral vector can be used (see Miller et al., Meth. Enzymol. 217:581-599 (1993)). These retroviral vectors contain the components necessary for the correct packaging of the viral genome and integration into the host cell DNA. The nucleic acid sequences encoding the antibody to be used in gene therapy are cloned into one or more vectors, which facilitates delivery of the gene into a patient. More detail about retroviral vectors can be found in Boesen et al., Biotherapy 6:291-302 (1994), which describes the use of a retroviral vector to deliver the *mdr1* gene to hematopoietic stem cells in order to make the stem cells more resistant to chemotherapy. Other references illustrating the use of retroviral vectors in gene therapy are: Clowes et al., J. Clin. Invest. 93:644-651 (1994); Kiem et al., Blood 83:1467-1473

administration to a subject of an expressed or expressible nucleic acid. In this embodiment of the invention, the nucleic acids produce their encoded protein that mediates a therapeutic effect.

[0278] Any of the methods for gene therapy available in the art can be used according to the present invention. Exemplary methods are described below.

[0279] For general reviews of the methods of gene therapy, see Goldspiel et al., *Clinical Pharmacy* 12:488-505 (1993); Wu and Wu, *Biotherapy* 3:87-95 (1991); Tolstoshev, *Ann. Rev. Pharmacol. Toxicol.* 32:573-596 (1993); Mulligan, *Science* 260:926-932 (1993); and Morgan and Anderson, *Ann. Rev. Biochem.* 62:191-217 (1993); May, *TIBTECH* 11(5):155-215 (1993). Methods commonly known in the art of recombinant DNA technology which can be used are described in Ausubel et al. (eds.), *Current Protocols in Molecular Biology*, John Wiley & Sons, NY (1993); and Kriegler, *Gene Transfer and Expression, A Laboratory Manual*, Stockton Press, NY (1990).

[0280] In a preferred embodiment, the compound comprises nucleic acid sequences encoding an antibody, said nucleic acid sequences being part of expression vectors that express the antibody or fragments or chimeric proteins or heavy or light chains thereof in a suitable host. In particular, such nucleic acid sequences have promoters operably linked to the antibody coding region, said promoter being inducible or constitutive, and, optionally, tissue-specific. In another particular embodiment, nucleic acid molecules are used in which the antibody coding sequences and any other desired sequences are flanked by regions that promote homologous recombination at a desired site in the genome, thus providing for intrachromosomal expression of the antibody encoding nucleic acids (Koller and Smithies, *Proc. Natl. Acad. Sci. USA* 86:8932-8935 (1989); Zijlstra et al., *Nature* 342:435-438 (1989). In specific embodiments, the expressed antibody molecule is a single chain antibody; alternatively, the nucleic acid sequences include sequences encoding both the heavy and light chains, or fragments thereof, of the antibody.

[0281] Delivery of the nucleic acids into a patient may be either direct, in which case the patient is directly exposed to the nucleic acid or nucleic acid-carrying vectors, or indirect, in which case, cells are first transformed with the nucleic acids *in vitro*, then transplanted into the patient. These two approaches are known, respectively, as *in vivo* or *ex vivo* gene therapy.

approaches are described in more detail below. Armed with the teachings provided herein, one of ordinary skill in the art will know how to use the antibodies of the present invention for diagnostic, monitoring or therapeutic purposes without undue experimentation.

[0274] The antibodies of this invention may be advantageously utilized in combination with other monoclonal or chimeric antibodies, or with lymphokines or hematopoietic growth factors (such as, e.g., IL-2, IL-3 and IL-7), for example, which serve to increase the number or activity of effector cells which interact with the antibodies.

[0275] The antibodies of the invention may be administered alone or in combination with other types of treatments (e.g., radiation therapy, chemotherapy, hormonal therapy, immunotherapy and anti-tumor agents). Generally, administration of products of a species origin or species reactivity (in the case of antibodies) that is the same species as that of the patient is preferred. Thus, in a preferred embodiment, human antibodies, fragments derivatives, analogs, or nucleic acids, are administered to a human patient for therapy or prophylaxis.

[0276] It is preferred to use high affinity and/or potent *in vivo* inhibiting and/or neutralizing antibodies against polypeptides or polynucleotides of the present invention, fragments or regions thereof, for both immunoassays directed to and therapy of disorders related to polynucleotides or polypeptides, including fragments thereof, of the present invention. Such antibodies, fragments, or regions, will preferably have an affinity for polynucleotides or polypeptides of the invention, including fragments thereof. Preferred binding affinities include those with a dissociation constant or K_d less than 5×10^{-2} M, 10^{-2} M, 5×10^{-3} M, 10^{-3} M, 5×10^{-4} M, 10^{-4} M, 5×10^{-5} M, 10^{-5} M, 5×10^{-6} M, 10^{-6} M, 5×10^{-7} M, 10^{-7} M, 5×10^{-8} M, 10^{-8} M, 5×10^{-9} M, 10^{-9} M, 5×10^{-10} M, 10^{-10} M, 5×10^{-11} M, 10^{-11} M, 5×10^{-12} M, 10^{-12} M, 5×10^{-13} M, 10^{-13} M, 5×10^{-14} M, 10^{-14} M, 5×10^{-15} M, and 10^{-15} M.

Gene Therapy

[0277] In a specific embodiment, nucleic acids comprising sequences encoding antibodies or functional derivatives thereof, are administered to treat, inhibit or prevent a disease or disorder associated with aberrant expression and/or activity of a polypeptide of the invention, by way of gene therapy. Gene therapy refers to therapy performed by the

diseases, disorders, or conditions described herein. The treatment and/or prevention of diseases, disorders, or conditions associated with aberrant expression and/or activity of a polypeptide of the invention includes, but is not limited to, alleviating symptoms associated with those diseases, disorders or conditions. Antibodies of the invention may be provided in pharmaceutically acceptable compositions as known in the art or as described herein.

[0272] In a specific and preferred embodiment, the present invention is directed to antibody-based therapies which involve administering antibodies of the invention to an animal, preferably a mammal, and most preferably a human, patient for treating one or more diseases, disorders, or conditions, including but not limited to: neural disorders, immune system disorders, muscular disorders, reproductive disorders, gastrointestinal disorders, pulmonary disorders, cardiovascular disorders, renal disorders, proliferative disorders, and/or cancerous diseases and conditions., and/or as described elsewhere herein. Therapeutic compounds of the invention include, but are not limited to, antibodies of the invention (e.g., antibodies directed to the full length protein expressed on the cell surface of a mammalian cell; antibodies directed to an epitope of a polypeptide of the invention (such as, for example, a predicted linear epitope shown in column 7 of Table 1; or a conformational epitope, including fragments, analogs and derivatives thereof as described herein) and nucleic acids encoding antibodies of the invention (including fragments, analogs and derivatives thereof and anti-idiotypic antibodies as described herein). The antibodies of the invention can be used to treat, inhibit or prevent diseases, disorders or conditions associated with aberrant expression and/or activity of a polypeptide of the invention, including, but not limited to, any one or more of the diseases, disorders, or conditions described herein. The treatment and/or prevention of diseases, disorders, or conditions associated with aberrant expression and/or activity of a polypeptide of the invention includes, but is not limited to, alleviating symptoms associated with those diseases, disorders or conditions. Antibodies of the invention may be provided in pharmaceutically acceptable compositions as known in the art or as described herein.

[0273] A summary of the ways in which the antibodies of the present invention may be used therapeutically includes binding polynucleotides or polypeptides of the present invention locally or systemically in the body or by direct cytotoxicity of the antibody, e.g. as mediated by complement (CDC) or by effector cells (ADCC). Some of these

knowledgeable as to the parameters that can be modified to increase the signal detected as well as other variations of ELISAs known in the art. For further discussion regarding ELISAs see, e.g., Ausubel et al, eds, (1994), Current Protocols in Molecular Biology, Vol. 1, John Wiley & Sons, Inc., New York, section 11.2.1.

[0269] The binding affinity of an antibody to an antigen and the off-rate of an antibody-antigen interaction can be determined by competitive binding assays. One example of a competitive binding assay is a radioimmunoassay comprising the incubation of labeled antigen (e.g., ^3H or ^{125}I) with the antibody of interest in the presence of increasing amounts of unlabeled antigen, and the detection of the antibody bound to the labeled antigen. The affinity of the antibody of interest for a particular antigen and the binding off-rates can be determined from the data by scatchard plot analysis. Competition with a second antibody can also be determined using radioimmunoassays. In this case, the antigen is incubated with antibody of interest conjugated to a labeled compound (e.g., ^3H or ^{125}I) in the presence of increasing amounts of an unlabeled second antibody.

[0270] Antibodies of the invention may be characterized using immunocytochemistry methods on cells (e.g., mammalian cells, such as CHO cells) transfected with a vector enabling the expression of an antigen or with vector alone using techniques commonly known in the art. Antibodies that bind antigen transfected cells, but not vector-only transfected cells, are antigen specific.

Therapeutic Uses

[0271] The present invention is further directed to antibody-based therapies which involve administering antibodies of the invention to an animal, preferably a mammal, and most preferably a human, patient for treating one or more of the disclosed diseases, disorders, or conditions. Therapeutic compounds of the invention include, but are not limited to, antibodies of the invention (including fragments, analogs and derivatives thereof as described herein) and nucleic acids encoding antibodies of the invention (including fragments, analogs and derivatives thereof and anti-idiotypic antibodies as described herein). The antibodies of the invention can be used to treat, inhibit or prevent diseases, disorders or conditions associated with aberrant expression and/or activity of a polypeptide of the invention, including, but not limited to, any one or more of the

western blot analysis. One of skill in the art would be knowledgeable as to the parameters that can be modified to increase the binding of the antibody to an antigen and decrease the background (e.g., pre-clearing the cell lysate with sepharose beads). For further discussion regarding immunoprecipitation protocols see, e.g., Ausubel et al., eds., (1994), *Current Protocols in Molecular Biology*, Vol. 1, John Wiley & Sons, Inc., New York, section 10.16.1.

[0267] Western blot analysis generally comprises preparing protein samples, electrophoresis of the protein samples in a polyacrylamide gel (e.g., 8%- 20% SDS-PAGE depending on the molecular weight of the antigen), transferring the protein sample from the polyacrylamide gel to a membrane such as nitrocellulose, PVDF or nylon, blocking the membrane in blocking solution (e.g., PBS with 3% BSA or non-fat milk), washing the membrane in washing buffer (e.g., PBS-Tween 20), blocking the membrane with primary antibody (the antibody of interest) diluted in blocking buffer, washing the membrane in washing buffer, blocking the membrane with a secondary antibody (which recognizes the primary antibody, e.g., an anti-human antibody) conjugated to an enzymatic substrate (e.g., horseradish peroxidase or alkaline phosphatase) or radioactive molecule (e.g., ^{32}P or ^{125}I) diluted in blocking buffer, washing the membrane in wash buffer, and detecting the presence of the antigen. One of skill in the art would be knowledgeable as to the parameters that can be modified to increase the signal detected and to reduce the background noise. For further discussion regarding western blot protocols see, e.g., Ausubel et al., eds., (1994), *Current Protocols in Molecular Biology*, Vol. 1, John Wiley & Sons, Inc., New York, section 10.8.1.

[0268] ELISAs comprise preparing antigen, coating the well of a 96 well microtiter plate with the antigen, adding the antibody of interest conjugated to a detectable compound such as an enzymatic substrate (e.g., horseradish peroxidase or alkaline phosphatase) to the well and incubating for a period of time, and detecting the presence of the antigen. In ELISAs the antibody of interest does not have to be conjugated to a detectable compound; instead, a second antibody (which recognizes the antibody of interest) conjugated to a detectable compound may be added to the well. Further, instead of coating the well with the antigen, the antibody may be coated to the well. In this case, a second antibody conjugated to a detectable compound may be added following the addition of the antigen of interest to the coated well. One of skill in the art would be

coated magnetic beads, "panning" with antibody attached to a solid matrix (i.e., plate), and flow cytometry (See, e.g., U.S. Patent 5,985,660; and Morrison *et al.*, *Cell*, 96:737-49 (1999)).

[0264] These techniques allow for the screening of particular populations of cells, such as might be found with hematological malignancies (i.e. minimal residual disease (MRD) in acute leukemic patients) and "non-self" cells in transplantations to prevent Graft-versus-Host Disease (GVHD). Alternatively, these techniques allow for the screening of hematopoietic stem and progenitor cells capable of undergoing proliferation and/or differentiation, as might be found in human umbilical cord blood.

Assays For Antibody Binding

[0265] The antibodies of the invention may be assayed for immunospecific binding by any method known in the art. The immunoassays which can be used include but are not limited to competitive and non-competitive assay systems using techniques such as western blots, radioimmunoassays, ELISA (enzyme linked immunosorbent assay), "sandwich" immunoassays, immunoprecipitation assays, precipitin reactions, gel diffusion precipitin reactions, immunodiffusion assays, agglutination assays, complement-fixation assays, immunoradiometric assays, fluorescent immunoassays, and protein A immunoassays, to name but a few. Such assays are routine and well known in the art (see, e.g., Ausubel *et al.*, eds, 1994, *Current Protocols in Molecular Biology*, Vol. 1, John Wiley & Sons, Inc., New York, which is incorporated by reference herein in its entirety). Exemplary immunoassays are described briefly below (but are not intended by way of limitation).

[0266] Immunoprecipitation protocols generally comprise lysing a population of cells in a lysis buffer such as RIPA buffer (1% NP-40 or Triton X- 100, 1% sodium deoxycholate, 0.1% SDS, 0.15 M NaCl, 0.01 M sodium phosphate at pH 7.2, 1% Trasylol) supplemented with protein phosphatase and/or protease inhibitors (e.g., EDTA, PMSF, aprotinin, sodium vanadate), adding the antibody of interest to the cell lysate, incubating for a period of time (e.g., 1-4 hours) at 4° C, adding protein A and/or protein G sepharose beads to the cell lysate, incubating for about an hour or more at 4° C, washing the beads in lysis buffer and resuspending the beads in SDS/sample buffer. The ability of the antibody of interest to immunoprecipitate a particular antigen can be assessed by, e.g.,

[0259] Antibodies may also be attached to solid supports, which are particularly useful for immunoassays or purification of the target antigen. Such solid supports include, but are not limited to, glass, cellulose, polyacrylamide, nylon, polystyrene, polyvinyl chloride or polypropylene.

[0260] Techniques for conjugating such therapeutic moiety to antibodies are well known. See, for example, Arnon et al., "Monoclonal Antibodies For Immunotargeting Of Drugs In Cancer Therapy", in *Monoclonal Antibodies And Cancer Therapy*, Reisfeld et al. (eds.), pp. 243-56 (Alan R. Liss, Inc. 1985); Hellstrom et al., "Antibodies For Drug Delivery", in *Controlled Drug Delivery* (2nd Ed.), Robinson et al. (eds.), pp. 623-53 (Marcel Dekker, Inc. 1987); Thorpe, "Antibody Carriers Of Cytotoxic Agents In Cancer Therapy: A Review", in *Monoclonal Antibodies '84: Biological And Clinical Applications*, Pinchera et al. (eds.), pp. 475-506 (1985); "Analysis, Results, And Future Prospective Of The Therapeutic Use Of Radiolabeled Antibody In Cancer Therapy", in *Monoclonal Antibodies For Cancer Detection And Therapy*, Baldwin et al. (eds.), pp. 303-16 (Academic Press 1985), and Thorpe et al., "The Preparation And Cytotoxic Properties Of Antibody-Toxin Conjugates", *Immunol. Rev.* 62:119-58 (1982).

[0261] Alternatively, an antibody can be conjugated to a second antibody to form an antibody heteroconjugate as described by Segal in U.S. Patent No. 4,676,980, which is incorporated herein by reference in its entirety.

[0262] An antibody, with or without a therapeutic moiety conjugated to it, administered alone or in combination with cytotoxic factor(s) and/or cytokine(s) can be used as a therapeutic.

Immunophenotyping

[0263] The antibodies of the invention may be utilized for immunophenotyping of cell lines and biological samples. Translation products of the gene of the present invention may be useful as cell-specific markers, or more specifically as cellular markers that are differentially expressed at various stages of differentiation and/or maturation of particular cell types. Monoclonal antibodies directed against a specific epitope, or combination of epitopes, will allow for the screening of cellular populations expressing the marker. Various techniques can be utilized using monoclonal antibodies to screen for cellular populations expressing the marker(s), and include magnetic separation using antibody-

[0257] Further, an antibody or fragment thereof may be conjugated to a therapeutic moiety such as a cytotoxin, e.g., a cytostatic or cytotoxic agent, a therapeutic agent or a radioactive metal ion, e.g., alpha-emitters such as, for example, ^{213}Bi . A cytotoxin or cytotoxic agent includes any agent that is detrimental to cells. Examples include paclitaxol, cytochalasin B, gramicidin D, ethidium bromide, emetine, mitomycin, etoposide, tenoposide, vincristine, vinblastine, colchicin, doxorubicin, daunorubicin, dihydroxy anthracin dione, mitoxantrone, mithramycin, actinomycin D, 1-dehydrotestosterone, glucocorticoids, procaine, tetracaine, lidocaine, propranolol, and puromycin and analogs or homologs thereof. Therapeutic agents include, but are not limited to, antimetabolites (e.g., methotrexate, 6-mercaptopurine, 6-thioguanine, cytarabine, 5-fluorouracil decarbazine), alkylating agents (e.g., mechlorethamine, thioepa chlorambucil, melphalan, carmustine (BSNU) and lomustine (CCNU), cyclophosphamide, busulfan, dibromomannitol, streptozotocin, mitomycin C, and cis- dichlorodiamine platinum (II) (DDP) cisplatin), anthracyclines (e.g., daunorubicin (formerly daunomycin) and doxorubicin), antibiotics (e.g., dactinomycin (formerly actinomycin), bleomycin, mithramycin, and anthramycin (AMC)), and anti-mitotic agents (e.g., vincristine and vinblastine).

[0258] The conjugates of the invention can be used for modifying a given biological response, the therapeutic agent or drug moiety is not to be construed as limited to classical chemical therapeutic agents. For example, the drug moiety may be a protein or polypeptide possessing a desired biological activity. Such proteins may include, for example, a toxin such as abrin, ricin A, pseudomonas exotoxin, or diphtheria toxin; a protein such as tumor necrosis factor, α -interferon, β -interferon, nerve growth factor, platelet derived growth factor, tissue plasminogen activator, an apoptotic agent, e.g., TNF- α , TNF- β , AIM I (See, International Publication No. WO 97/33899), AIM II (See, International Publication No. WO 97/34911), Fas Ligand (Takahashi *et al.*, *Int. Immunol.*, 6:1567-1574 (1994)), VEGI (See, International Publication No. WO 99/23105), a thrombotic agent or an anti- angiogenic agent, e.g., angiostatin or endostatin; or, biological response modifiers such as, for example, lymphokines, interleukin-1 ("IL-1"), interleukin-2 ("IL-2"), interleukin-6 ("IL-6"), granulocyte macrophage colony stimulating factor ("GM-CSF"), granulocyte colony stimulating factor ("G-CSF"), or other growth factors.

Molecular Recognition 8:52-58 (1995); Johanson et al., J. Biol. Chem. 270:9459-9471 (1995)).

[0255] Moreover, the antibodies or fragments thereof of the present invention can be fused to marker sequences, such as a peptide to facilitate purification. In preferred embodiments, the marker amino acid sequence is a hexa-histidine peptide, such as the tag provided in a pQE vector (QIAGEN, Inc., 9259 Eton Avenue, Chatsworth, CA, 91311), among others, many of which are commercially available. As described in Gentz et al., Proc. Natl. Acad. Sci. USA 86:821-824 (1989), for instance, hexa-histidine provides for convenient purification of the fusion protein. Other peptide tags useful for purification include, but are not limited to, the "HA" tag, which corresponds to an epitope derived from the influenza hemagglutinin protein (Wilson et al., Cell 37:767 (1984)) and the "flag" tag.

[0256] The present invention further encompasses antibodies or fragments thereof conjugated to a diagnostic or therapeutic agent. The antibodies can be used diagnostically to, for example, monitor the development or progression of a tumor as part of a clinical testing procedure to, e.g., determine the efficacy of a given treatment regimen. Detection can be facilitated by coupling the antibody to a detectable substance. Examples of detectable substances include various enzymes, prosthetic groups, fluorescent materials, luminescent materials, bioluminescent materials, radioactive materials, positron emitting metals using various positron emission tomographies, and nonradioactive paramagnetic metal ions. The detectable substance may be coupled or conjugated either directly to the antibody (or fragment thereof) or indirectly, through an intermediate (such as, for example, a linker known in the art) using techniques known in the art. See, for example, U.S. Patent No. 4,741,900 for metal ions which can be conjugated to antibodies for use as diagnostics according to the present invention. Examples of suitable enzymes include horseradish peroxidase, alkaline phosphatase, beta-galactosidase, or acetylcholinesterase; examples of suitable prosthetic group complexes include streptavidin/biotin and avidin/biotin; examples of suitable fluorescent materials include umbelliferone, fluorescein, fluorescein isothiocyanate, rhodamine, dichlorotriazinylamine fluorescein, dansyl chloride or phycoerythrin; an example of a luminescent material includes luminol; examples of bioluminescent materials include luciferase, luciferin, and aequorin; and examples of suitable radioactive material include ¹²⁵I, ¹³¹I, ¹¹¹In or ⁹⁹Tc.

CH1 domain, CH2 domain, and CH3 domain or any combination of whole domains or portions thereof. The polypeptides may also be fused or conjugated to the above antibody portions to form multimers. For example, Fc portions fused to the polypeptides of the present invention can form dimers through disulfide bonding between the Fc portions. Higher multimeric forms can be made by fusing the polypeptides to portions of IgA and IgM. Methods for fusing or conjugating the polypeptides of the present invention to antibody portions are known in the art. See, e.g., U.S. Patent Nos. 5,336,603; 5,622,929; 5,359,046; 5,349,053; 5,447,851; 5,112,946; EP 307,434; EP 367,166; PCT publications WO 96/04388; WO 91/06570; Ashkenazi et al., Proc. Natl. Acad. Sci. USA 88:10535-10539 (1991); Zheng et al., J. Immunol. 154:5590-5600 (1995); and Vil et al., Proc. Natl. Acad. Sci. USA 89:11337-11341 (1992) (said references incorporated by reference in their entireties).

[0254] As discussed, *supra*, the polypeptides corresponding to a polypeptide, polypeptide fragment, or a variant of SEQ ID NO:Y may be fused or conjugated to the above antibody portions to increase the *in vivo* half life of the polypeptides or for use in immunoassays using methods known in the art. Further, the polypeptides corresponding to SEQ ID NO:Y may be fused or conjugated to the above antibody portions to facilitate purification. One reported example describes chimeric proteins consisting of the first two domains of the human CD4-polypeptide and various domains of the constant regions of the heavy or light chains of mammalian immunoglobulins. See EP 394,827; and Traunecker et al., Nature 331:84-86 (1988). The polypeptides of the present invention fused or conjugated to an antibody having disulfide-linked dimeric structures (due to the IgG) may also be more efficient in binding and neutralizing other molecules, than the monomeric secreted protein or protein fragment alone. See, for example, Fountoulakis et al., J. Biochem. 270:3958-3964 (1995). In many cases, the Fc part in a fusion protein is beneficial in therapy and diagnosis, and thus can result in, for example, improved pharmacokinetic properties. See, for example, EP A 232,262. Alternatively, deleting the Fc part after the fusion protein has been expressed, detected, and purified, would be desired. For example, the Fc portion may hinder therapy and diagnosis if the fusion protein is used as an antigen for immunizations. In drug discovery, for example, human proteins, such as hIL-5, have been fused with Fc portions for the purpose of high-throughput screening assays to identify antagonists of hIL-5. (See, Bennett et al., J.

Nature 322:52 (1986); Kohler, Proc. Natl. Acad. Sci. USA 77:2197 (1980)). The coding sequences for the heavy and light chains may comprise cDNA or genomic DNA.

[0251] Once an antibody molecule of the invention has been produced by an animal, chemically synthesized, or recombinantly expressed, it may be purified by any method known in the art for purification of an immunoglobulin molecule, for example, by chromatography (e.g., ion exchange, affinity, particularly by affinity for the specific antigen after Protein A, and sizing column chromatography), centrifugation, differential solubility, or by any other standard technique for the purification of proteins. In addition, the antibodies of the present invention or fragments thereof can be fused to heterologous polypeptide sequences described herein or otherwise known in the art, to facilitate purification.

[0252] The present invention encompasses antibodies recombinantly fused or chemically conjugated (including both covalently and non-covalently conjugations) to a polypeptide (or portion thereof, preferably at least 10, 20, 30, 40, 50, 60, 70, 80, 90 or 100 amino acids of the polypeptide) of the present invention to generate fusion proteins. The fusion does not necessarily need to be direct, but may occur through linker sequences. The antibodies may be specific for antigens other than polypeptides (or portion thereof, preferably at least 10, 20, 30, 40, 50, 60, 70, 80, 90 or 100 amino acids of the polypeptide) of the present invention. For example, antibodies may be used to target the polypeptides of the present invention to particular cell types, either *in vitro* or *in vivo*, by fusing or conjugating the polypeptides of the present invention to antibodies specific for particular cell surface receptors. Antibodies fused or conjugated to the polypeptides of the present invention may also be used in *in vitro* immunoassays and purification methods using methods known in the art. See e.g., Harbor et al., *supra*, and PCT publication WO 93/21232; EP 439,095; Naramura et al., Immunol. Lett. 39:91-99 (1994); U.S. Patent 5,474,981; Gillies et al., PNAS 89:1428-1432 (1992); Fell et al., J. Immunol. 146:2446-2452 (1991), which are incorporated by reference in their entireties.

[0253] The present invention further includes compositions comprising the polypeptides of the present invention fused or conjugated to antibody domains other than the variable regions. For example, the polypeptides of the present invention may be fused or conjugated to an antibody Fc region, or portion thereof. The antibody portion fused to a polypeptide of the present invention may comprise the constant region, hinge region,

[0248] The expression levels of an antibody molecule can be increased by vector amplification (for a review, see Bebbington and Hentschel, The use of vectors based on gene amplification for the expression of cloned genes in mammalian cells in DNA cloning, Vol.3. (Academic Press, New York, 1987)). When a marker in the vector system expressing antibody is amplifiable, increase in the level of inhibitor present in culture of host cell will increase the number of copies of the marker gene. Since the amplified region is associated with the antibody gene, production of the antibody will also increase (Crouse et al., Mol. Cell. Biol. 3:257 (1983)).

[0249] Vectors which use glutamine synthase (GS) or DHFR as the selectable markers can be amplified in the presence of the drugs methionine sulfoximine or methotrexate, respectively. An advantage of glutamine synthase based vectors are the availability of cell lines (e.g., the murine myeloma cell line, NS0) which are glutamine synthase negative. Glutamine synthase expression systems can also function in glutamine synthase expressing cells (e.g. Chinese Hamster Ovary (CHO) cells) by providing additional inhibitor to prevent the functioning of the endogenous gene. A glutamine synthase expression system and components thereof are detailed in PCT publications: WO87/04462; WO86/05807; WO89/01036; WO89/10404; and WO91/06657 which are incorporated in their entireties by reference herein. Additionally, glutamine synthase expression vectors that may be used according to the present invention are commercially available from suppliers, including, for example Lonza Biologics, Inc. (Portsmouth, NH). Expression and production of monoclonal antibodies using a GS expression system in murine myeloma cells is described in Bebbington *et al.*, *Bio/technology* 10:169(1992) and in Biblia and Robinson *Biotechnol. Prog.* 11:1 (1995) which are incorporated in their entireties by reference herein.

[0250] The host cell may be co-transfected with two expression vectors of the invention, the first vector encoding a heavy chain derived polypeptide and the second vector encoding a light chain derived polypeptide. The two vectors may contain identical selectable markers which enable equal expression of heavy and light chain polypeptides. Alternatively, a single vector may be used which encodes, and is capable of expressing, both heavy and light chain polypeptides. In such situations, the light chain should be placed before the heavy chain to avoid an excess of toxic free heavy chain (Proudfoot,

replication, host cells can be transformed with DNA controlled by appropriate expression control elements (e.g., promoter, enhancer, sequences, transcription terminators, polyadenylation sites, etc.), and a selectable marker. Following the introduction of the foreign DNA, engineered cells may be allowed to grow for 1-2 days in an enriched media, and then are switched to a selective media. The selectable marker in the recombinant plasmid confers resistance to the selection and allows cells to stably integrate the plasmid into their chromosomes and grow to form foci which in turn can be cloned and expanded into cell lines. This method may advantageously be used to engineer cell lines which express the antibody molecule. Such engineered cell lines may be particularly useful in screening and evaluation of compounds that interact directly or indirectly with the antibody molecule.

[0247] A number of selection systems may be used, including but not limited to the herpes simplex virus thymidine kinase (Wigler et al., *Cell* 11:223 (1977)), hypoxanthine-guanine phosphoribosyltransferase (Szybalska & Szybalski, *Proc. Natl. Acad. Sci. USA* 48:202 (1992)), and adenine phosphoribosyltransferase (Lowy et al., *Cell* 22:817 (1980)) genes can be employed in tk-, hgp^{rt}- or ap^{rt}- cells, respectively. Also, antimetabolite resistance can be used as the basis of selection for the following genes: dhfr, which confers resistance to methotrexate (Wigler et al., *Natl. Acad. Sci. USA* 77:357 (1980); O'Hare et al., *Proc. Natl. Acad. Sci. USA* 78:1527 (1981)); gpt, which confers resistance to mycophenolic acid (Mulligan & Berg, *Proc. Natl. Acad. Sci. USA* 78:2072 (1981)); neo, which confers resistance to the aminoglycoside G-418 (*Clinical Pharmacology* 12:488-505; Wu and Wu, *Biotherapy* 3:87-95 (1991); Tolstoshev, *Ann. Rev. Pharmacol. Toxicol.* 32:573-596 (1993); Mulligan, *Science* 260:926-932 (1993); and Morgan and Anderson, *Ann. Rev. Biochem.* 62:191-217 (1993); May, 1993, *TIB TECH* 11(5):155-215 (1993)); and hyg^r, which confers resistance to hygromycin (Santerre et al., *Gene* 30:147 (1984)). Methods commonly known in the art of recombinant DNA technology may be routinely applied to select the desired recombinant clone, and such methods are described, for example, in Ausubel et al. (eds.), *Current Protocols in Molecular Biology*, John Wiley & Sons, NY (1993); Kriegler, *Gene Transfer and Expression, A Laboratory Manual*, Stockton Press, NY (1990); and in Chapters 12 and 13, Dracopoli et al. (eds.), *Current Protocols in Human Genetics*, John Wiley & Sons, NY (1994); Colberre-Garapin et al., *J. Mol. Biol.* 150:1 (1981), which are incorporated by reference herein in their entireties.

[0244] In mammalian host cells, a number of viral-based expression systems may be utilized. In cases where an adenovirus is used as an expression vector, the antibody coding sequence of interest may be ligated to an adenovirus transcription/translation control complex, e.g., the late promoter and tripartite leader sequence. This chimeric gene may then be inserted in the adenovirus genome by *in vitro* or *in vivo* recombination. Insertion in a non-essential region of the viral genome (e.g., region E1 or E3) will result in a recombinant virus that is viable and capable of expressing the antibody molecule in infected hosts. (e.g., see Logan & Shenk, Proc. Natl. Acad. Sci. USA 81:355-359 (1984)). Specific initiation signals may also be required for efficient translation of inserted antibody coding sequences. These signals include the ATG initiation codon and adjacent sequences. Furthermore, the initiation codon must be in phase with the reading frame of the desired coding sequence to ensure translation of the entire insert. These exogenous translational control signals and initiation codons can be of a variety of origins, both natural and synthetic. The efficiency of expression may be enhanced by the inclusion of appropriate transcription enhancer elements, transcription terminators, etc. (see Bittner et al., Methods in Enzymol. 153:51-544 (1987)).

[0245] In addition, a host cell strain may be chosen which modulates the expression of the inserted sequences, or modifies and processes the gene product in the specific fashion desired. Such modifications (e.g., glycosylation) and processing (e.g., cleavage) of protein products may be important for the function of the protein. Different host cells have characteristic and specific mechanisms for the post-translational processing and modification of proteins and gene products. Appropriate cell lines or host systems can be chosen to ensure the correct modification and processing of the foreign protein expressed. To this end, eukaryotic host cells which possess the cellular machinery for proper processing of the primary transcript, glycosylation, and phosphorylation of the gene product may be used. Such mammalian host cells include but are not limited to CHO, VERY, BHK, Hela, COS, MDCK, 293, 3T3, WI38, and in particular, breast cancer cell lines such as, for example, BT483, Hs578T, HTB2, BT20 and T47D, and normal mammary gland cell line such as, for example, CRL7030 and Hs578Bst.

[0246] For long-term, high-yield production of recombinant proteins, stable expression is preferred. For example, cell lines which stably express the antibody molecule may be engineered. Rather than using expression vectors which contain viral origins of

CHO, BHK, 293, 3T3 cells) harboring recombinant expression constructs containing promoters derived from the genome of mammalian cells (e.g., metallothionein promoter) or from mammalian viruses (e.g., the adenovirus late promoter; the vaccinia virus 7.5K promoter). Preferably, bacterial cells such as *Escherichia coli*, and more preferably, eukaryotic cells, especially for the expression of whole recombinant antibody molecule, are used for the expression of a recombinant antibody molecule. For example, mammalian cells such as Chinese hamster ovary cells (CHO), in conjunction with a vector such as the major intermediate early gene promoter element from human cytomegalovirus is an effective expression system for antibodies (Foecking et al., *Gene* 45:101 (1986); Cockett et al., *Bio/Technology* 8:2 (1990)).

[0242] In bacterial systems, a number of expression vectors may be advantageously selected depending upon the use intended for the antibody molecule being expressed. For example, when a large quantity of such a protein is to be produced, for the generation of pharmaceutical compositions of an antibody molecule, vectors which direct the expression of high levels of fusion protein products that are readily purified may be desirable. Such vectors include, but are not limited, to the *E. coli* expression vector pUR278 (Ruther et al., *EMBO J.* 2:1791 (1983)), in which the antibody coding sequence may be ligated individually into the vector in frame with the lac Z coding region so that a fusion protein is produced; pIN vectors (Inouye & Inouye, *Nucleic Acids Res.* 13:3101-3109 (1985); Van Heeke & Schuster, *J. Biol. Chem.* 24:5503-5509 (1989)); and the like. pGEX vectors may also be used to express foreign polypeptides as fusion proteins with glutathione S-transferase (GST). In general, such fusion proteins are soluble and can easily be purified from lysed cells by adsorption and binding to matrix glutathione-agarose beads followed by elution in the presence of free glutathione. The pGEX vectors are designed to include thrombin or factor Xa protease cleavage sites so that the cloned target gene product can be released from the GST moiety.

[0243] In an insect system, *Autographa californica* nuclear polyhedrosis virus (AcNPV) is used as a vector to express foreign genes. The virus grows in *Spodoptera frugiperda* cells. The antibody coding sequence may be cloned individually into non-essential regions (for example the polyhedrin gene) of the virus and placed under control of an AcNPV promoter (for example the polyhedrin promoter).

can be used to construct expression vectors containing antibody coding sequences and appropriate transcriptional and translational control signals. These methods include, for example, *in vitro* recombinant DNA techniques, synthetic techniques, and *in vivo* genetic recombination. The invention, thus, provides replicable vectors comprising a nucleotide sequence encoding an antibody molecule of the invention, or a heavy or light chain thereof, or a heavy or light chain variable domain, operably linked to a promoter. Such vectors may include the nucleotide sequence encoding the constant region of the antibody molecule (see, e.g., PCT Publication WO 86/05807; PCT Publication WO 89/01036; and U.S. Patent No. 5,122,464) and the variable domain of the antibody may be cloned into such a vector for expression of the entire heavy or light chain.

[0240] The expression vector is transferred to a host cell by conventional techniques and the transfected cells are then cultured by conventional techniques to produce an antibody of the invention. Thus, the invention includes host cells containing a polynucleotide encoding an antibody of the invention, or a heavy or light chain thereof, or a single chain antibody of the invention, operably linked to a heterologous promoter. In preferred embodiments for the expression of double-chained antibodies, vectors encoding both the heavy and light chains may be co-expressed in the host cell for expression of the entire immunoglobulin molecule, as detailed below.

[0241] A variety of host-expression vector systems may be utilized to express the antibody molecules of the invention. Such host-expression systems represent vehicles by which the coding sequences of interest may be produced and subsequently purified, but also represent cells which may, when transformed or transfected with the appropriate nucleotide coding sequences, express an antibody molecule of the invention *in situ*. These include but are not limited to microorganisms such as bacteria (e.g., *E. coli*, *B. subtilis*) transformed with recombinant bacteriophage DNA, plasmid DNA or cosmid DNA expression vectors containing antibody coding sequences; yeast (e.g., *Saccharomyces*, *Pichia*) transformed with recombinant yeast expression vectors containing antibody coding sequences; insect cell systems infected with recombinant virus expression vectors (e.g., baculovirus) containing antibody coding sequences; plant cell systems infected with recombinant virus expression vectors (e.g., cauliflower mosaic virus, CaMV; tobacco mosaic virus, TMV) or transformed with recombinant plasmid expression vectors (e.g., Ti plasmid) containing antibody coding sequences; or mammalian cell systems (e.g., COS,

[0236] In addition, techniques developed for the production of "chimeric antibodies" (Morrison et al., Proc. Natl. Acad. Sci. 81:851-855 (1984); Neuberger et al., Nature 312:604-608 (1984); Takeda et al., Nature 314:452-454 (1985)) by splicing genes from a mouse antibody molecule of appropriate antigen specificity together with genes from a human antibody molecule of appropriate biological activity can be used. As described *supra*, a chimeric antibody is a molecule in which different portions are derived from different animal species, such as those having a variable region derived from a murine mAb and a human immunoglobulin constant region, e.g., humanized antibodies.

[0237] Alternatively, techniques described for the production of single chain antibodies (U.S. Patent No. 4,946,778; Bird, Science 242:423- 42 (1988); Huston et al., Proc. Natl. Acad. Sci. USA 85:5879-5883 (1988); and Ward et al., Nature 334:544-54 (1989)) can be adapted to produce single chain antibodies. Single chain antibodies are formed by linking the heavy and light chain fragments of the Fv region via an amino acid bridge, resulting in a single chain polypeptide. Techniques for the assembly of functional Fv fragments in *E. coli* may also be used (Skerra et al., Science 242:1038- 1041 (1988)).

Methods of Producing Antibodies

[0238] The antibodies of the invention can be produced by any method known in the art for the synthesis of antibodies, in particular, by chemical synthesis or preferably, by recombinant expression techniques. Methods of producing antibodies include, but are not limited to, hybridoma technology, EBV transformation, and other methods discussed herein as well as through the use recombinant DNA technology, as discussed below.

[0239] Recombinant expression of an antibody of the invention, or fragment, derivative or analog thereof, (e.g., a heavy or light chain of an antibody of the invention or a single chain antibody of the invention), requires construction of an expression vector containing a polynucleotide that encodes the antibody. Once a polynucleotide encoding an antibody molecule or a heavy or light chain of an antibody, or portion thereof (preferably containing the heavy or light chain variable domain), of the invention has been obtained, the vector for the production of the antibody molecule may be produced by recombinant DNA technology using techniques well known in the art. Thus, methods for preparing a protein by expressing a polynucleotide containing an antibody encoding nucleotide sequence are described herein. Methods which are well known to those skilled in the art

Amplified nucleic acids generated by PCR may then be cloned into replicable cloning vectors using any method well known in the art.

[0234] Once the nucleotide sequence and corresponding amino acid sequence of the antibody is determined, the nucleotide sequence of the antibody may be manipulated using methods well known in the art for the manipulation of nucleotide sequences, e.g., recombinant DNA techniques, site directed mutagenesis, PCR, etc. (see, for example, the techniques described in Sambrook et al., 1990, *Molecular Cloning, A Laboratory Manual*, 2d Ed., Cold Spring Harbor Laboratory, Cold Spring Harbor, NY and Ausubel et al., eds., 1998, *Current Protocols in Molecular Biology*, John Wiley & Sons, NY, which are both incorporated by reference herein in their entireties), to generate antibodies having a different amino acid sequence, for example to create amino acid substitutions, deletions, and/or insertions.

[0235] In a specific embodiment, the amino acid sequence of the heavy and/or light chain variable domains may be inspected to identify the sequences of the complementarity determining regions (CDRs) by methods that are well known in the art, e.g., by comparison to known amino acid sequences of other heavy and light chain variable regions to determine the regions of sequence hypervariability. Using routine recombinant DNA techniques, one or more of the CDRs may be inserted within framework regions, e.g., into human framework regions to humanize a non-human antibody, as described *supra*. The framework regions may be naturally occurring or consensus framework regions, and preferably human framework regions (see, e.g., Chothia et al., *J. Mol. Biol.* 278: 457-479 (1998) for a listing of human framework regions). Preferably, the polynucleotide generated by the combination of the framework regions and CDRs encodes an antibody that specifically binds a polypeptide of the invention. Preferably, as discussed *supra*, one or more amino acid substitutions may be made within the framework regions, and, preferably, the amino acid substitutions improve binding of the antibody to its antigen. Additionally, such methods may be used to make amino acid substitutions or deletions of one or more variable region cysteine residues participating in an intrachain disulfide bond to generate antibody molecules lacking one or more intrachain disulfide bonds. Other alterations to the polynucleotide are encompassed by the present invention and within the skill of the art.

al., J. Mol. Biol. 291:1129-1134 (1999); Wirtz and Steipe, Protein Sci. 8:2245-2250 (1999); Zhu et al., J. Immunol. Methods 231:207-222 (1999); and references cited therein.

Polynucleotides Encoding Antibodies

[0231] The invention further provides polynucleotides comprising a nucleotide sequence encoding an antibody of the invention and fragments thereof. The invention also encompasses polynucleotides that hybridize under stringent or alternatively, under lower stringency hybridization conditions, e.g., as defined *supra*, to polynucleotides that encode an antibody, preferably, that specifically binds to a polypeptide of the invention, preferably, an antibody that binds to a polypeptide having the amino acid sequence of SEQ ID NO:Y, to a polypeptide encoded by a portion of SEQ ID NO:X as defined in columns 8 and 9 of Table 2, and/or to a polypeptide encoded by the cDNA contained in Clone ID NO:Z.

[0232] The polynucleotides may be obtained, and the nucleotide sequence of the polynucleotides determined, by any method known in the art. For example, if the nucleotide sequence of the antibody is known, a polynucleotide encoding the antibody may be assembled from chemically synthesized oligonucleotides (e.g., as described in Kutmeier et al., BioTechniques 17:242 (1994)), which, briefly, involves the synthesis of overlapping oligonucleotides containing portions of the sequence encoding the antibody, annealing and ligating of those oligonucleotides, and then amplification of the ligated oligonucleotides by PCR.

[0233] Alternatively, a polynucleotide encoding an antibody may be generated from nucleic acid from a suitable source. If a clone containing a nucleic acid encoding a particular antibody is not available, but the sequence of the antibody molecule is known, a nucleic acid encoding the immunoglobulin may be chemically synthesized or obtained from a suitable source (e.g., an antibody cDNA library, or a cDNA library generated from, or nucleic acid, preferably poly A+ RNA, isolated from, any tissue or cells expressing the antibody, such as hybridoma cells selected to express an antibody of the invention) by PCR amplification using synthetic primers hybridizable to the 3' and 5' ends of the sequence or by cloning using an oligonucleotide probe specific for the particular gene sequence to identify, e.g., a cDNA clone from a cDNA library that encodes the antibody.

human antibodies directed against a selected antigen using technology similar to that described above.

[0228] Completely human antibodies which recognize a selected epitope can be generated using a technique referred to as "guided selection." In this approach a selected non-human monoclonal antibody, e.g., a mouse antibody, is used to guide the selection of a completely human antibody recognizing the same epitope. (Jespers et al., *Bio/technology* 12:899-903 (1988)).

[0229] Further, antibodies to the polypeptides of the invention can, in turn, be utilized to generate anti-idiotypic antibodies that "mimic" polypeptides of the invention using techniques well known to those skilled in the art. (See, e.g., Greenspan & Bona, *FASEB J.* 7(5):437-444; (1989) and Nissinoff, J. *Immunol.* 147(8):2429-2438 (1991)). For example, antibodies which bind to and competitively inhibit polypeptide multimerization and/or binding of a polypeptide of the invention to a ligand can be used to generate anti-idiotypes that "mimic" the polypeptide multimerization and/or binding domain and, as a consequence, bind to and neutralize polypeptide and/or its ligand. Such neutralizing anti-idiotypes or Fab fragments of such anti-idiotypes can be used in therapeutic regimens to neutralize polypeptide ligand(s)/receptor(s). For example, such anti-idiotypic antibodies can be used to bind a polypeptide of the invention and/or to bind its ligand(s)/receptor(s), and thereby block its biological activity. Alternatively, antibodies which bind to and enhance polypeptide multimerization and/or binding, and/or receptor/ligand multimerization, binding and/or signaling can be used to generate anti-idiotypes that function as agonists of a polypeptide of the invention and/or its ligand/receptor. Such agonistic anti-idiotypes or Fab fragments of such anti-idiotypes can be used in therapeutic regimens as agonists of the polypeptides of the invention or its ligand(s)/receptor(s). For example, such anti-idiotypic antibodies can be used to bind a polypeptide of the invention and/or to bind its ligand(s)/receptor(s), and thereby promote or enhance its biological activity.

[0230] Intrabodies of the invention can be produced using methods known in the art, such as those disclosed and reviewed in Chen et al., *Hum. Gene Ther.* 5:595-601 (1994); Marasco, W.A., *Gene Ther.* 4:11-15 (1997); Rondon and Marasco, *Annu. Rev. Microbiol.* 51:257-283 (1997); Proba et al., *J. Mol. Biol.* 275:245-253 (1998); Cohen et al., *Oncogene* 17:2445-2456 (1998); Ohage and Steipe, *J. Mol. Biol.* 291:1119-1128 (1999); Ohage et

libraries derived from human immunoglobulin sequences. See also, U.S. Patent Nos. 4,444,887 and 4,716,111; and PCT publications WO 98/46645, WO 98/50433, WO 98/24893, WO 98/16654, WO 96/34096, WO 96/33735, and WO 91/10741; each of which is incorporated herein by reference in its entirety.

[0227] Human antibodies can also be produced using transgenic mice which are incapable of expressing functional endogenous immunoglobulins, but which can express human immunoglobulin genes. For example, the human heavy and light chain immunoglobulin gene complexes may be introduced randomly or by homologous recombination into mouse embryonic stem cells. Alternatively, the human variable region, constant region, and diversity region may be introduced into mouse embryonic stem cells in addition to the human heavy and light chain genes. The mouse heavy and light chain immunoglobulin genes may be rendered non-functional separately or simultaneously with the introduction of human immunoglobulin loci by homologous recombination. In particular, homozygous deletion of the JH region prevents endogenous antibody production. The modified embryonic stem cells are expanded and microinjected into blastocysts to produce chimeric mice. The chimeric mice are then bred to produce homozygous offspring which express human antibodies. The transgenic mice are immunized in the normal fashion with a selected antigen, e.g., all or a portion of a polypeptide of the invention. Monoclonal antibodies directed against the antigen can be obtained from the immunized, transgenic mice using conventional hybridoma technology. The human immunoglobulin transgenes harbored by the transgenic mice rearrange during B cell differentiation, and subsequently undergo class switching and somatic mutation. Thus, using such a technique, it is possible to produce therapeutically useful IgG, IgA, IgM and IgE antibodies. For an overview of this technology for producing human antibodies, see Lonberg and Huszar, *Int. Rev. Immunol.* 13:65-93 (1995). For a detailed discussion of this technology for producing human antibodies and human monoclonal antibodies and protocols for producing such antibodies, see, e.g., PCT publications WO 98/24893; WO 92/01047; WO 96/34096; WO 96/33735; European Patent No. 0 598 877; U.S. Patent Nos. 5,413,923; 5,625,126; 5,633,425; 5,569,825; 5,661,016; 5,545,806; 5,814,318; 5,885,793; 5,916,771; 5,939,598; 6,075,181; and 6,114,598, which are incorporated by reference herein in their entirety. In addition, companies such as Abgenix, Inc. (Freemont, CA) and Genpharm (San Jose, CA) can be engaged to provide

[0225] Examples of techniques which can be used to produce single-chain Fvs and antibodies include those described in U.S. Patents 4,946,778 and 5,258,498; Huston et al., *Methods in Enzymology* 203:46-88 (1991); Shu et al., *PNAS* 90:7995-7999 (1993); and Skerra et al., *Science* 240:1038-1040 (1988). For some uses, including *in vivo* use of antibodies in humans and *in vitro* detection assays, it may be preferable to use chimeric, humanized, or human antibodies. A chimeric antibody is a molecule in which different portions of the antibody are derived from different animal species, such as antibodies having a variable region derived from a murine monoclonal antibody and a human immunoglobulin constant region. Methods for producing chimeric antibodies are known in the art. See e.g., Morrison, *Science* 229:1202 (1985); Oi et al., *BioTechniques* 4:214 (1986); Gillies et al., (1989) *J. Immunol. Methods* 125:191-202; U.S. Patent Nos. 5,807,715; 4,816,567; and 4,816,397, which are incorporated herein by reference in their entirety. Humanized antibodies are antibody molecules from non-human species antibody that binds the desired antigen having one or more complementarity determining regions (CDRs) from the non-human species and a framework regions from a human immunoglobulin molecule. Often, framework residues in the human framework regions will be substituted with the corresponding residue from the CDR donor antibody to alter, preferably improve, antigen binding. These framework substitutions are identified by methods well known in the art, e.g., by modeling of the interactions of the CDR and framework residues to identify framework residues important for antigen binding and sequence comparison to identify unusual framework residues at particular positions. (See, e.g., Queen et al., U.S. Patent No. 5,585,089; Riechmann et al., *Nature* 332:323 (1988), which are incorporated herein by reference in their entirety.) Antibodies can be humanized using a variety of techniques known in the art including, for example, CDR-grafting (EP 239,400; PCT publication WO 91/09967; U.S. Patent Nos. 5,225,539; 5,530,101; and 5,585,089), veneering or resurfacing (EP 592,106; EP 519,596; Padlan, *Molecular Immunology* 28(4/5):489-498 (1991); Studnicka et al., *Protein Engineering* 7(6):805-814 (1994); Roguska et al., *PNAS* 91:969-973 (1994)), and chain shuffling (U.S. Patent No. 5,565,332).

[0226] Completely human antibodies are particularly desirable for therapeutic treatment of human patients. Human antibodies can be made by a variety of methods known in the art including phage display methods described above using antibody

fragments contain the variable region, the light chain constant region and the CH1 domain of the heavy chain.

[0223] For example, the antibodies of the present invention can also be generated using various phage display methods known in the art. In phage display methods, functional antibody domains are displayed on the surface of phage particles which carry the polynucleotide sequences encoding them. In a particular embodiment, such phage can be utilized to display antigen binding domains expressed from a repertoire or combinatorial antibody library (e.g., human or murine). Phage expressing an antigen binding domain that binds the antigen of interest can be selected or identified with antigen, e.g., using labeled antigen or antigen bound or captured to a solid surface or bead. Phage used in these methods are typically filamentous phage including fd and M13 binding domains expressed from phage with Fab, Fv or disulfide stabilized Fv antibody domains recombinantly fused to either the phage gene III or gene VIII protein. Examples of phage display methods that can be used to make the antibodies of the present invention include those disclosed in Brinkman et al., *J. Immunol. Methods* 182:41-50 (1995); Ames et al., *J. Immunol. Methods* 184:177-186 (1995); Kettleborough et al., *Eur. J. Immunol.* 24:952-958 (1994); Persic et al., *Gene* 187 9-18 (1997); Burton et al., *Advances in Immunology* 57:191-280 (1994); PCT application No. PCT/GB91/01134; PCT publications WO 90/02809; WO 91/10737; WO 92/01047; WO 92/18619; WO 93/11236; WO 95/15982; WO 95/20401; and U.S. Patent Nos. 5,698,426; 5,223,409; 5,403,484; 5,580,717; 5,427,908; 5,750,753; 5,821,047; 5,571,698; 5,427,908; 5,516,637; 5,780,225; 5,658,727; 5,733,743 and 5,969,108; each of which is incorporated herein by reference in its entirety.

[0224] As described in the above references, after phage selection, the antibody coding regions from the phage can be isolated and used to generate whole antibodies, including human antibodies, or any other desired antigen binding fragment, and expressed in any desired host, including mammalian cells, insect cells, plant cells, yeast, and bacteria, e.g., as described in detail below. For example, techniques to recombinantly produce Fab, Fab' and F(ab')₂ fragments can also be employed using methods known in the art such as those disclosed in PCT publication WO 92/22324; Mullinax et al., *BioTechniques* 12(6):864-869 (1992); and Sawai et al., *AJRI* 34:26-34 (1995); and Better et al., *Science* 240:1041-1043 (1988) (said references incorporated by reference in their entireties).

generating EBV-transformed B cell lines are commonly known in the art, such as, for example, the protocol outlined in Chapter 7.22 of *Current Protocols in Immunology*, Coligan et al., Eds., 1994, John Wiley & Sons, NY, which is hereby incorporated in its entirety by reference. The source of B cells for transformation is commonly human peripheral blood, but B cells for transformation may also be derived from other sources including, but not limited to, lymph nodes, tonsil, spleen, tumor tissue, and infected tissues. Tissues are generally made into single cell suspensions prior to EBV transformation. Additionally, steps may be taken to either physically remove or inactivate T cells (e.g., by treatment with cyclosporin A) in B cell-containing samples, because T cells from individuals seropositive for anti-EBV antibodies can suppress B cell immortalization by EBV.

[0221] In general, the sample containing human B cells is inoculated with EBV, and cultured for 3-4 weeks. A typical source of EBV is the culture supernatant of the B95-8 cell line (ATCC #VR-1492). Physical signs of EBV transformation can generally be seen towards the end of the 3-4 week culture period. By phase-contrast microscopy, transformed cells may appear large, clear, hairy and tend to aggregate in tight clusters of cells. Initially, EBV lines are generally polyclonal. However, over prolonged periods of cell cultures, EBV lines may become monoclonal or polyclonal as a result of the selective outgrowth of particular B cell clones. Alternatively, polyclonal EBV transformed lines may be subcloned (e.g., by limiting dilution culture) or fused with a suitable fusion partner and plated at limiting dilution to obtain monoclonal B cell lines. Suitable fusion partners for EBV transformed cell lines include mouse myeloma cell lines (e.g., SP2/0, X63-Ag8.653), heteromyeloma cell lines (human x mouse; e.g., SPAM-8, SBC-H20, and CB-F7), and human cell lines (e.g., GM 1500, SKO-007, RPMI 8226, and KR-4). Thus, the present invention also provides a method of generating polyclonal or monoclonal human antibodies against polypeptides of the invention or fragments thereof, comprising EBV-transformation of human B cells.

[0222] Antibody fragments which recognize specific epitopes may be generated by known techniques. For example, Fab and F(ab')₂ fragments of the invention may be produced by proteolytic cleavage of immunoglobulin molecules, using enzymes such as papain (to produce Fab fragments) or pepsin (to produce F(ab')₂ fragments). F(ab')₂

[0217] Monoclonal antibodies can be prepared using a wide variety of techniques known in the art including the use of hybridoma, recombinant, and phage display technologies, or a combination thereof. For example, monoclonal antibodies can be produced using hybridoma techniques including those known in the art and taught, for example, in Harlow et al., *Antibodies: A Laboratory Manual*, (Cold Spring Harbor Laboratory Press, 2nd ed. 1988); Hammerling, et al., in: *Monoclonal Antibodies and T-Cell Hybridomas* 563-681 (Elsevier, N.Y., 1981) (said references incorporated by reference in their entirety). The term "monoclonal antibody" as used herein is not limited to antibodies produced through hybridoma technology. The term "monoclonal antibody" refers to an antibody that is derived from a single clone, including any eukaryotic, prokaryotic, or phage clone, and not the method by which it is produced.

[0218] Methods for producing and screening for specific antibodies using hybridoma technology are routine and well known in the art and are discussed in detail in the Examples. In a non-limiting example, mice can be immunized with a polypeptide of the invention or a cell expressing such peptide. Once an immune response is detected, e.g., antibodies specific for the antigen are detected in the mouse serum, the mouse spleen is harvested and splenocytes isolated. The splenocytes are then fused by well known techniques to any suitable myeloma cells, for example cells from cell line SP20 available from the ATCC. Hybridomas are selected and cloned by limited dilution. The hybridoma clones are then assayed by methods known in the art for cells that secrete antibodies capable of binding a polypeptide of the invention. Ascites fluid, which generally contains high levels of antibodies, can be generated by immunizing mice with positive hybridoma clones.

[0219] Accordingly, the present invention provides methods of generating monoclonal antibodies as well as antibodies produced by the method comprising culturing a hybridoma cell secreting an antibody of the invention wherein, preferably, the hybridoma is generated by fusing splenocytes isolated from a mouse immunized with an antigen of the invention with myeloma cells and then screening the hybridomas resulting from the fusion for hybridoma clones that secrete an antibody able to bind a polypeptide of the invention.

[0220] Another well known method for producing both polyclonal and monoclonal human B cell lines is transformation using Epstein Barr Virus (EBV). Protocols for

[0214] As discussed in more detail below, the antibodies of the present invention may be used either alone or in combination with other compositions. The antibodies may further be recombinantly fused to a heterologous polypeptide at the N- or C-terminus or chemically conjugated (including covalent and non-covalent conjugations) to polypeptides or other compositions. For example, antibodies of the present invention may be recombinantly fused or conjugated to molecules useful as labels in detection assays and effector molecules such as heterologous polypeptides, drugs, radionuclides, or toxins. See, e.g., PCT publications WO 92/08495; WO 91/14438; WO 89/12624; U.S. Patent No. 5,314,995; and EP 396,387; the disclosures of which are incorporated herein by reference in their entireties.

[0215] The antibodies of the invention include derivatives that are modified, i.e., by the covalent attachment of any type of molecule to the antibody such that covalent attachment does not prevent the antibody from generating an anti-idiotypic response. For example, but not by way of limitation, the antibody derivatives include antibodies that have been modified, e.g., by glycosylation, acetylation, pegylation, phosphorylation, amidation, derivatization by known protecting/blocking groups, proteolytic cleavage, linkage to a cellular ligand or other protein, etc. Any of numerous chemical modifications may be carried out by known techniques, including, but not limited to specific chemical cleavage, acetylation, formylation, metabolic synthesis of tunicamycin, etc. Additionally, the derivative may contain one or more non-classical amino acids.

[0216] The antibodies of the present invention may be generated by any suitable method known in the art. Polyclonal antibodies to an antigen-of-interest can be produced by various procedures well known in the art. For example, a polypeptide of the invention can be administered to various host animals including, but not limited to, rabbits, mice, rats, etc. to induce the production of sera containing polyclonal antibodies specific for the antigen. Various adjuvants may be used to increase the immunological response, depending on the host species, and include but are not limited to, Freund's (complete and incomplete), mineral gels such as aluminum hydroxide, surface active substances such as lysolecithin, pluronic polyols, polyanions, peptides, oil emulsions, keyhole limpet hemocyanins, dinitrophenol, and potentially useful human adjuvants such as BCG (bacille Calmette-Guerin) and corynebacterium parvum. Such adjuvants are also well known in the art.

[0912] In yet an additional embodiment, the Therapeutics of the invention are delivered by way of a pump (*see* Langer, *supra*; Sefton, CRC Crit. Ref. Biomed. Eng. 14:201 (1987); Buchwald et al., Surgery 88:507 (1980); Saudek et al., N. Engl. J. Med. 321:574 (1989)).

[0913] Other controlled release systems are discussed in the review by Langer (*Science* 249:1527-1533 (1990)).

[0914] For parenteral administration, in one embodiment, the Therapeutic is formulated generally by mixing it at the desired degree of purity, in a unit dosage injectable form (solution, suspension, or emulsion), with a pharmaceutically acceptable carrier, i.e., one that is non-toxic to recipients at the dosages and concentrations employed and is compatible with other ingredients of the formulation. For example, the formulation preferably does not include oxidizing agents and other compounds that are known to be deleterious to the Therapeutic.

[0915] Generally, the formulations are prepared by contacting the Therapeutic uniformly and intimately with liquid carriers or finely divided solid carriers or both. Then, if necessary, the product is shaped into the desired formulation. Preferably the carrier is a parenteral carrier, more preferably a solution that is isotonic with the blood of the recipient. Examples of such carrier vehicles include water, saline, Ringer's solution, and dextrose solution. Non-aqueous vehicles such as fixed oils and ethyl oleate are also useful herein, as well as liposomes.

[0916] The carrier suitably contains minor amounts of additives such as substances that enhance isotonicity and chemical stability. Such materials are non-toxic to recipients at the dosages and concentrations employed, and include buffers such as phosphate, citrate, succinate, acetic acid, and other organic acids or their salts; antioxidants such as ascorbic acid; low molecular weight (less than about ten residues) polypeptides, e.g., polyarginine or tripeptides; proteins, such as serum albumin, gelatin, or immunoglobulins; hydrophilic polymers such as polyvinylpyrrolidone; amino acids, such as glycine, glutamic acid, aspartic acid, or arginine; monosaccharides, disaccharides, and other carbohydrates including cellulose or its derivatives, glucose, manose, or dextrans; chelating agents such as EDTA; sugar alcohols such as mannitol or sorbitol; counterions such as sodium; and/or nonionic surfactants such as polysorbates, poloxamers, or PEG.

[0917] The Therapeutic is typically formulated in such vehicles at a concentration of about 0.1 mg/ml to 100 mg/ml, preferably 1-10 mg/ml, at a pH of about 3 to 8. It will be understood that the use of certain of the foregoing excipients, carriers, or stabilizers will result in the formation of polypeptide salts.

[0918] Any pharmaceutical used for therapeutic administration can be sterile. Sterility is readily accomplished by filtration through sterile filtration membranes (e.g., 0.2 micron membranes). Therapeutics generally are placed into a container having a sterile access port, for example, an intravenous solution bag or vial having a stopper pierceable by a hypodermic injection needle.

[0919] Therapeutics ordinarily will be stored in unit or multi-dose containers, for example, sealed ampoules or vials, as an aqueous solution or as a lyophilized formulation for reconstitution. As an example of a lyophilized formulation, 10-ml vials are filled with 5 ml of sterile-filtered 1% (w/v) aqueous Therapeutic solution, and the resulting mixture is lyophilized. The infusion solution is prepared by reconstituting the lyophilized Therapeutic using bacteriostatic Water-for-Injection.

[0920] The invention also provides a pharmaceutical pack or kit comprising one or more containers filled with one or more of the ingredients of the Therapeutics of the invention. Associated with such container(s) can be a notice in the form prescribed by a governmental agency regulating the manufacture, use or sale of pharmaceuticals or biological products, which notice reflects approval by the agency of manufacture, use or sale for human administration. In addition, the Therapeutics may be employed in conjunction with other therapeutic compounds.

[0921] The Therapeutics of the invention may be administered alone or in combination with adjuvants. Adjuvants that may be administered with the Therapeutics of the invention include, but are not limited to, alum, alum plus deoxycholate (ImmunoAg), MTP-PE (Biocine Corp.), QS21 (Genentech, Inc.), BCG (e.g., THERACYS®), MPL and nonviable preparations of *Corynebacterium parvum*. In a specific embodiment, Therapeutics of the invention are administered in combination with alum. In another specific embodiment, Therapeutics of the invention are administered in combination with QS-21. Further adjuvants that may be administered with the Therapeutics of the invention include, but are not limited to, Monophosphoryl lipid immunomodulator, AdjuVax 100a, QS-21, QS-18, CRL1005, Aluminum salts, MF-59, and Virosomal adjuvant technology.

Vaccines that may be administered with the Therapeutics of the invention include, but are not limited to, vaccines directed toward protection against MMR (measles, mumps, rubella), polio, varicella, tetanus/diphtheria, hepatitis A, hepatitis B, haemophilus influenzae B, whooping cough, pneumonia, influenza, Lyme's Disease, rotavirus, cholera, yellow fever, Japanese encephalitis, poliomyelitis, rabies, typhoid fever, and pertussis. Combinations may be administered either concomitantly, e.g., as an admixture, separately but simultaneously or concurrently; or sequentially. This includes presentations in which the combined agents are administered together as a therapeutic mixture, and also procedures in which the combined agents are administered separately but simultaneously, e.g., as through separate intravenous lines into the same individual. Administration "in combination" further includes the separate administration of one of the compounds or agents given first, followed by the second.

[0922] The Therapeutics of the invention may be administered alone or in combination with other therapeutic agents. Therapeutic agents that may be administered in combination with the Therapeutics of the invention, include but not limited to, chemotherapeutic agents, antibiotics, steroidal and non-steroidal anti-inflammatories, conventional immunotherapeutic agents, and/or therapeutic treatments described below. Combinations may be administered either concomitantly, e.g., as an admixture, separately but simultaneously or concurrently; or sequentially. This includes presentations in which the combined agents are administered together as a therapeutic mixture, and also procedures in which the combined agents are administered separately but simultaneously, e.g., as through separate intravenous lines into the same individual. Administration "in combination" further includes the separate administration of one of the compounds or agents given first, followed by the second.

[0923] In certain embodiments, the Therapeutics of the invention are administered in combination with an anticoagulant. Anticoagulants that may be administered with the compositions of the invention include, but are not limited to, heparin, low molecular weight heparin, warfarin sodium (e.g., COUMADIN®), dicumarol, 4-hydroxycoumarin, anisindione (e.g., MIRADON™), acenocoumarol (e.g., nicoumalone, SINTHROME™), indan-1,3-dione, phenprocoumon (e.g., MARCUMAR™), ethyl biscoumacetate (e.g., TROMEXAN™), and aspirin. In a specific embodiment, compositions of the invention are administered in combination with heparin and/or warfarin. In another specific

embodiment, compositions of the invention are administered in combination with warfarin. In another specific embodiment, compositions of the invention are administered in combination with warfarin and aspirin. In another specific embodiment, compositions of the invention are administered in combination with heparin. In another specific embodiment, compositions of the invention are administered in combination with heparin and aspirin.

[0924] In another embodiment, the Therapeutics of the invention are administered in combination with thrombolytic drugs. Thrombolytic drugs that may be administered with the compositions of the invention include, but are not limited to, plasminogen, lys-plasminogen, alpha2-antiplasmin, streptokinae (e.g., KABIKINASE™), antirespace (e.g., EMINASE™), tissue plasminogen activator (t-PA, altevase, ACTIVASE™), urokinase (e.g., ABBOKINASE™), sauruplase, (Prourokinase, single chain urokinase), and aminocaproic acid (e.g., AMICAR™). In a specific embodiment, compositions of the invention are administered in combination with tissue plasminogen activator and aspirin.

[0925] In another embodiment, the Therapeutics of the invention are administered in combination with antiplatelet drugs. Antiplatelet drugs that may be administered with the compositions of the invention include, but are not limited to, aspirin, dipyridamole (e.g., PERSANTINE™), and ticlopidine (e.g., TICLID™).

[0926] In specific embodiments, the use of anti-coagulants, thrombolytic and/or antiplatelet drugs in combination with Therapeutics of the invention is contemplated for the prevention, diagnosis, and/or treatment of thrombosis, arterial thrombosis, venous thrombosis, thromboembolism, pulmonary embolism, atherosclerosis, myocardial infarction, transient ischemic attack, unstable angina. In specific embodiments, the use of anticoagulants, thrombolytic drugs and/or antiplatelet drugs in combination with Therapeutics of the invention is contemplated for the prevention of occlusion of saphenous grafts, for reducing the risk of periprocedural thrombosis as might accompany angioplasty procedures, for reducing the risk of stroke in patients with atrial fibrillation including nonrheumatic atrial fibrillation, for reducing the risk of embolism associated with mechanical heart valves and or mitral valves disease. Other uses for the therapeutics of the invention, alone or in combination with antiplatelet, anticoagulant, and/or thrombolytic drugs, include, but are not limited to, the prevention of occlusions in extracorporeal devices (e.g., intravascular canulas, vascular access shunts in hemodialysis

patients, hemodialysis machines, and cardiopulmonary bypass machines).

[0927] In certain embodiments, Therapeutics of the invention are administered in combination with antiretroviral agents, nucleoside/nucleotide reverse transcriptase inhibitors (NRTIs), non-nucleoside reverse transcriptase inhibitors (NNRTIs), and/or protease inhibitors (PIs). NRTIs that may be administered in combination with the Therapeutics of the invention, include, but are not limited to, RETROVIR™ (zidovudine/AZT), VIDEX™ (didanosine/ddI), HIVID™ (zalcitabine/ddC), ZERIT™ (stavudine/d4T), EPIVIR™ (lamivudine/3TC), and COMBIVIR™ (zidovudine/lamivudine). NNRTIs that may be administered in combination with the Therapeutics of the invention, include, but are not limited to, VIRAMUNE™ (nevirapine), RESCRIPTOR™ (delavirdine), and SUSTIVA™ (efavirenz). Protease inhibitors that may be administered in combination with the Therapeutics of the invention, include, but are not limited to, CRIVAN™ (indinavir), NORVIR™ (ritonavir), INVIRASE™ (saquinavir), and VIRACEPT™ (nelfinavir). In a specific embodiment, antiretroviral agents, nucleoside reverse transcriptase inhibitors, non-nucleoside reverse transcriptase inhibitors, and/or protease inhibitors may be used in any combination with Therapeutics of the invention to treat AIDS and/or to prevent or treat HIV infection.

[0928] Additional NRTIs include LODENOSINE™ (F-ddA; an acid-stable adenosine NRTI; Triangle/Abbott); COVIRACIL™ (emtricitabine/FTC; structurally related to lamivudine (3TC) but with 3- to 10-fold greater activity *in vitro*; Triangle/Abbott); dOTC (BCH-10652, also structurally related to lamivudine but retains activity against a substantial proportion of lamivudine-resistant isolates; Biochem Pharma); Adefovir (refused approval for anti-HIV therapy by FDA; Gilead Sciences); PREVEON® (Adefovir Dipivoxil, the active prodrug of adefovir; its active form is PMEA-pp); TENOFOVIR™ (bis-POC PMPA, a PMPA prodrug; Gilead); DAPD/DXG (active metabolite of DAPD; Triangle/Abbott); D-D4FC (related to 3TC, with activity against AZT/3TC-resistant virus); GW420867X (Glaxo Wellcome); ZIAGEN™ (abacavir/159U89; Glaxo Wellcome Inc.); CS-87 (3'-azido-2',3'-dideoxyuridine; WO 99/66936); and S-acyl-2-thioethyl (SATE)-bearing prodrug forms of β -L-FD4C and β -L-FddC (WO 98/17281).

[0929] Additional NNRTIs include COACTINON™ (Emivirine/MKC-442, potent NNRTI of the HEPT class; Triangle/Abbott); CAPRAVIRINE™ (AG-1549/S-1153, a next generation NNRTI with activity against viruses containing the K103N mutation; Agouron); PNU-142721 (has 20- to 50-fold greater activity than its predecessor delavirdine and is active against K103N mutants; Pharmacia & Upjohn); DPC-961 and DPC-963 (second-generation derivatives of efavirenz, designed to be active against viruses with the K103N mutation; DuPont); GW-420867X (has 25-fold greater activity than HBY097 and is active against K103N mutants; Glaxo Wellcome); CALANOLIDE A (naturally occurring agent from the latex tree; active against viruses containing either or both the Y181C and K103N mutations); and Propolis (WO 99/49830).

[0930] Additional protease inhibitors include LOPINAVIR™ (ABT378/r; Abbott Laboratories); BMS-232632 (an azapeptide; Bristol-Myers Squibb); TIPRANAVIR™ (PNU-140690, a non-peptic dihydropyrone; Pharmacia & Upjohn); PD-178390 (a nonpeptidic dihydropyrone; Parke-Davis); BMS 232632 (an azapeptide; Bristol-Myers Squibb); L-756,423 (an indinavir analog; Merck); DMP-450 (a cyclic urea compound; Avid & DuPont); AG-1776 (a peptidomimetic with *in vitro* activity against protease inhibitor-resistant viruses; Agouron); VX-175/GW-433908 (phosphate prodrug of amprenavir; Vertex & Glaxo Wellcome); CGP61755 (Ciba); and AGENERASE™ (amprenavir; Glaxo Wellcome Inc.).

[0931] Additional antiretroviral agents include fusion inhibitors/gp41 binders. Fusion inhibitors/gp41 binders include T-20 (a peptide from residues 643-678 of the HIV gp41 transmembrane protein ectodomain which binds to gp41 in its resting state and prevents transformation to the fusogenic state; Trimeris) and T-1249 (a second-generation fusion inhibitor; Trimeris).

[0932] Additional antiretroviral agents include fusion inhibitors/chemokine receptor antagonists. Fusion inhibitors/chemokine receptor antagonists include CXCR4 antagonists such as AMD 3100 (a bicyclam), SDF-1 and its analogs, and ALX40-4C (a cationic peptide), T22 (an 18 amino acid peptide; Trimeris) and the T22 analogs T134 and T140; CCR5 antagonists such as RANTES (9-68), AOP-RANTES, NNY-RANTES, and TAK-779; and CCR5/CXCR4 antagonists such as NSC 651016 (a distamycin analog). Also included are CCR2B, CCR3, and CCR6 antagonists. Chemokine receptor agonists such as RANTES, SDF-1, MIP-1 α , MIP-1 β , etc., may also inhibit fusion.

[0933] Additional antiretroviral agents include integrase inhibitors. Integrase inhibitors include dicaffeoylquinic (DFQA) acids; L-chicoric acid (a dicaffeoyltartaric (DCTA) acid); quinalizarin (QLC) and related anthraquinones; ZINTEVIR™ (AR 177, an oligonucleotide that probably acts at cell surface rather than being a true integrase inhibitor; Arondex); and naphthols such as those disclosed in WO 98/50347.

[0934] Additional antiretroviral agents include hydroxyurea-like compounds such as BCX-34 (a purine nucleoside phosphorylase inhibitor; Biocryst); ribonucleotide reductase inhibitors such as DIDOX™ (Molecules for Health); inosine monophosphate dehydrogenase (IMPDH) inhibitors such as VX-497 (Vertex); and mycopholic acids such as CellCept (mycophenolate mofetil; Roche).

[0935] Additional antiretroviral agents include inhibitors of viral integrase, inhibitors of viral genome nuclear translocation such as arylene bis(methylketone) compounds; inhibitors of HIV entry such as AOP-RANTES, NNY-RANTES, RANTES-IgG fusion protein, soluble complexes of RANTES and glycosaminoglycans (GAG), and AMD-3100; nucleocapsid zinc finger inhibitors such as dithiane compounds; targets of HIV Tat and Rev; and pharmacoenhancers such as ABT-378.

[0936] Other antiretroviral therapies and adjunct therapies include cytokines and lymphokines such as MIP-1 α , MIP-1 β , SDF-1 α , IL-2, PROLEUKIN™ (aldesleukin/L2-7001; Chiron), IL-4, IL-10, IL-12, and IL-13; interferons such as IFN- α 2a; antagonists of TNFs, NF κ B, GM-CSF, M-CSF, and IL-10; agents that modulate immune activation such as cyclosporin and prednisone; vaccines such as Remune™ (HIV Immunogen), APL 400-003 (Apollon), recombinant gp120 and fragments, bivalent (B/E) recombinant envelope glycoprotein, rgp120CM235, MN rgp120, SF-2 rgp120, gp120/soluble CD4 complex, Delta JR-FL protein, branched synthetic peptide derived from discontinuous gp120 C3/C4 domain, fusion-competent immunogens, and Gag, Pol, Nef, and Tat vaccines; gene-based therapies such as genetic suppressor elements (GSEs; WO 98/54366), and intrakines (genetically modified CC chemokines targetted to the ER to block surface expression of newly synthesized CCR5 (Yang *et al.*, *PNAS* 94:11567-72 (1997); Chen *et al.*, *Nat. Med.* 3:1110-16 (1997)); antibodies such as the anti-CXCR4 antibody 12G5, the anti-CCR5 antibodies 2D7, 5C7, PA8, PA9, PA10, PA11, PA12, and PA14, the anti-CD4 antibodies Q4120 and RPA-T4, the anti-CCR3 antibody 7B11, the anti-gp120 antibodies 17b, 48d, 447-52D, 257-D, 268-D and 50.1, anti-Tat antibodies, anti-TNF- α antibodies,

and monoclonal antibody 33A; aryl hydrocarbon (AH) receptor agonists and antagonists such as TCDD, 3,3',4,4',5-pentachlorobiphenyl, 3,3',4,4'-tetrachlorobiphenyl, and α -naphthoflavone (WO 98/30213); and antioxidants such as γ -L-glutamyl-L-cysteine ethyl ester (γ -GCE; WO 99/56764).

[0937] In a further embodiment, the Therapeutics of the invention are administered in combination with an antiviral agent. Antiviral agents that may be administered with the Therapeutics of the invention include, but are not limited to, acyclovir, ribavirin, amantadine, and remantidine.

[0938] In other embodiments, Therapeutics of the invention may be administered in combination with anti-opportunistic infection agents. Anti-opportunistic agents that may be administered in combination with the Therapeutics of the invention, include, but are not limited to, TRIMETHOPRIM-SULFAMETHOXAZOLE™, DAPSONE™, PENTAMIDINE™, ATOVAQUONE™, ISONIAZID™, RIFAMPIN™, PYRAZINAMIDE™, ETHAMBUTOL™, RIFABUTIN™, CLARITHROMYCIN™, AZITHROMYCIN™, GANCICLOVIR™, FOSCARNET™, CIDOFOVIR™, FLUCONAZOLE™, ITRACONAZOLE™, KETOCONAZOLE™, ACYCLOVIR™, FAMCICOLVIR™, PYRIMETHAMINE™, LEUCOVORIN™, NEUPOGEN™ (filgrastim/G-CSF), and LEUKINE™ (sargramostim/GM-CSF). In a specific embodiment, Therapeutics of the invention are used in any combination with TRIMETHOPRIM-SULFAMETHOXAZOLE™, DAPSONE™, PENTAMIDINE™, and/or ATOVAQUONE™ to prophylactically treat or prevent an opportunistic *Pneumocystis carinii* pneumonia infection. In another specific embodiment, Therapeutics of the invention are used in any combination with ISONIAZID™, RIFAMPIN™, PYRAZINAMIDE™, and/or ETHAMBUTOL™ to prophylactically treat or prevent an opportunistic *Mycobacterium avium* complex infection. In another specific embodiment, Therapeutics of the invention are used in any combination with RIFABUTIN™, CLARITHROMYCIN™, and/or AZITHROMYCIN™ to prophylactically treat or prevent an opportunistic *Mycobacterium tuberculosis* infection. In another specific embodiment, Therapeutics of the invention are used in any combination with GANCICLOVIR™, FOSCARNET™, and/or CIDOFOVIR™ to prophylactically treat or prevent an opportunistic cytomegalovirus infection. In another specific embodiment, Therapeutics

of the invention are used in any combination with FLUCONAZOLE™, ITRACONAZOLE™, and/or KETOCONAZOLE™ to prophylactically treat or prevent an opportunistic fungal infection. In another specific embodiment, Therapeutics of the invention are used in any combination with ACYCLOVIR™ and/or FAMCICOLVIR™ to prophylactically treat or prevent an opportunistic herpes simplex virus type I and/or type II infection. In another specific embodiment, Therapeutics of the invention are used in any combination with PYRIMETHAMINE™ and/or LEUCOVORIN™ to prophylactically treat or prevent an opportunistic *Toxoplasma gondii* infection. In another specific embodiment, Therapeutics of the invention are used in any combination with LEUCOVORIN™ and/or NEUPOGEN™ to prophylactically treat or prevent an opportunistic bacterial infection.

[0939] In a further embodiment, the Therapeutics of the invention are administered in combination with an antibiotic agent. Antibiotic agents that may be administered with the Therapeutics of the invention include, but are not limited to, amoxicillin, beta-lactamases, aminoglycosides, beta-lactam (glycopeptide), beta-lactamases, Clindamycin, chloramphenicol, cephalosporins, ciprofloxacin, erythromycin, fluoroquinolones, macrolides, metronidazole, penicillins, quinolones, rapamycin, rifampin, streptomycin, sulfonamide, tetracyclines, trimethoprim, trimethoprim-sulfamethoxazole, and vancomycin.

[0940] In other embodiments, the Therapeutics of the invention are administered in combination with immunestimulants. Immunostimulants that may be administered in combination with the Therapeutics of the invention include, but are not limited to, levamisole (e.g., ERGAMISOL™), isoprinosine (e.g. INOSIPLEX™), interferons (e.g. interferon alpha), and interleukins (e.g., IL-2).

[0941] In other embodiments, Therapeutics of the invention are administered in combination with immunosuppressive agents. Immunosuppressive agents that may be administered in combination with the Therapeutics of the invention include, but are not limited to, steroids, cyclosporine, cyclosporine analogs, cyclophosphamide methylprednisone, prednisone, azathioprine, FK-506, 15-deoxyspergualin, and other immunosuppressive agents that act by suppressing the function of responding T cells. Other immunosuppressive agents that may be administered in combination with the Therapeutics of the invention include, but are not limited to, prednisolone, methotrexate,

thalidomide, methoxsalen, rapamycin, leflunomide, mizoribine (BREDININ™), brequinar, deoxyspergualin, and azaspirane (SKF 105685), ORTHOCLONE OKT® 3 (muromonab-CD3), SANDIMMUNE™, NEORAL™, SANGDYA™ (cyclosporine), PROGRAF® (FK506, tacrolimus), CELLCEPT® (mycophenolate mofetil, of which the active metabolite is mycophenolic acid), IMURAN™ (azathioprine), glucocorticosteroids, adrenocortical steroids such as DELTASONE™ (prednisone) and HYDELTRASOL™ (prednisolone), FOLEX™ and MEXATE™ (methotrexate), OXSORALEN-ULTRA™ (methoxsalen) and RAPAMUNE™ (sirolimus). In a specific embodiment, immunosuppressants may be used to prevent rejection of organ or bone marrow transplantation.

[0942] In an additional embodiment, Therapeutics of the invention are administered alone or in combination with one or more intravenous immune globulin preparations. Intravenous immune globulin preparations that may be administered with the Therapeutics of the invention include, but not limited to, GAMMAR™, IVEEGAM™, SANDOGLOBULIN™, GAMMAGARD S/D™, ATGAM™ (antithymocyte globulin), and GAMIMUNE™. In a specific embodiment, Therapeutics of the invention are administered in combination with intravenous immune globulin preparations in transplantation therapy (e.g., bone marrow transplant).

[0943] In certain embodiments, the Therapeutics of the invention are administered alone or in combination with an anti-inflammatory agent. Anti-inflammatory agents that may be administered with the Therapeutics of the invention include, but are not limited to, corticosteroids (e.g. betamethasone, budesonide, cortisone, dexamethasone, hydrocortisone, methylprednisolone, prednisolone, prednisone, and triamcinolone), nonsteroidal anti-inflammatory drugs (e.g., diclofenac, diflunisal, etodolac, fenoprofen, floctafenine, flurbiprofen, ibuprofen, indomethacin, ketoprofen, meclofenamate, mefenamic acid, meloxicam, nabumetone, naproxen, oxaprozin, phenylbutazone, piroxicam, sulindac, tenoxicam, tiaprofenic acid, and tolmetin.), as well as antihistamines, aminoarylcarboxylic acid derivatives, arylacetic acid derivatives, arylbutyric acid derivatives, arylcarboxylic acids, arylpropionic acid derivatives, pyrazoles, pyrazolones, salicylic acid derivatives, thiazinecarboxamides, e-acetamidocaproic acid, S-adenosylmethionine, 3-amino-4-hydroxybutyric acid, amixetrine, bendazac, benzydamine, bucolome, difenpiramide, ditazol, emorfazone, guaiazulene, nabumetone,

nimesulide, orgotein, oxaceprol, paranyline, perisoxal, pifoxime, proquazone, proxazole, and tenidap.

[0944] In an additional embodiment, the compositions of the invention are administered alone or in combination with an anti-angiogenic agent. Anti-angiogenic agents that may be administered with the compositions of the invention include, but are not limited to, Angiostatin (Entremed, Rockville, MD), Troponin-1 (Boston Life Sciences, Boston, MA), anti-Invasive Factor, retinoic acid and derivatives thereof, paclitaxel (Taxol), Suramin, Tissue Inhibitor of Metalloproteinase-1, Tissue Inhibitor of Metalloproteinase-2, VEGI, Plasminogen Activator Inhibitor-1, Plasminogen Activator Inhibitor-2, and various forms of the lighter "d group" transition metals.

[0945] Lighter "d group" transition metals include, for example, vanadium, molybdenum, tungsten, titanium, niobium, and tantalum species. Such transition metal species may form transition metal complexes. Suitable complexes of the above-mentioned transition metal species include oxo transition metal complexes.

[0946] Representative examples of vanadium complexes include oxo vanadium complexes such as vanadate and vanadyl complexes. Suitable vanadate complexes include metavanadate and orthovanadate complexes such as, for example, ammonium metavanadate, sodium metavanadate, and sodium orthovanadate. Suitable vanadyl complexes include, for example, vanadyl acetylacetonate and vanadyl sulfate including vanadyl sulfate hydrates such as vanadyl sulfate mono- and trihydrates.

[0947] Representative examples of tungsten and molybdenum complexes also include oxo complexes. Suitable oxo tungsten complexes include tungstate and tungsten oxide complexes. Suitable tungstate complexes include ammonium tungstate, calcium tungstate, sodium tungstate dihydrate, and tungstic acid. Suitable tungsten oxides include tungsten (IV) oxide and tungsten (VI) oxide. Suitable oxo molybdenum complexes include molybdate, molybdenum oxide, and molybdenyl complexes. Suitable molybdate complexes include ammonium molybdate and its hydrates, sodium molybdate and its hydrates, and potassium molybdate and its hydrates. Suitable molybdenum oxides include molybdenum (VI) oxide, molybdenum (VI) oxide, and molybdic acid. Suitable molybdenyl complexes include, for example, molybdenyl acetylacetonate. Other suitable tungsten and molybdenum complexes include hydroxo derivatives derived from, for example, glycerol, tartaric acid, and sugars.

[0948] A wide variety of other anti-angiogenic factors may also be utilized within the context of the present invention. Representative examples include, but are not limited to, platelet factor 4; protamine sulphate; sulphated chitin derivatives (prepared from queen crab shells), (Murata et al., *Cancer Res.* 51:22-26, (1991)); Sulphated Polysaccharide Peptidoglycan Complex (SP- PG) (the function of this compound may be enhanced by the presence of steroids such as estrogen, and tamoxifen citrate); Staurosporine; modulators of matrix metabolism, including for example, proline analogs, cishydroxyproline, d,L-3,4-dehydroproline, Thiaproline, alpha,alpha-dipyridyl, aminopropionitrile fumarate; 4-propyl-5-(4-pyridinyl)-2(3H)-oxazolone; Methotrexate; Mitoxantrone; Heparin; Interferons; 2 Macroglobulin-serum; ChIMP-3 (Pavloff et al., *J. Bio. Chem.* 267:17321-17326, (1992)); Chymostatin (Tomkinson et al., *Biochem J.* 286:475-480, (1992)); Cyclodextrin Tetradecasulfate; Eponemycin; Camptothecin; Fumagillin (Ingber et al., *Nature* 348:555-557, (1990)); Gold Sodium Thiomalate ("GST"; Matsubara and Ziff, *J. Clin. Invest.* 79:1440-1446, (1987)); anticollagenase-serum; alpha2-antiplasmin (Holmes et al., *J. Biol. Chem.* 262(4):1659-1664, (1987)); Bisantrone (National Cancer Institute); Lobenzarit disodium (N-(2)-carboxyphenyl-4- chloroanthronilic acid disodium or "CCA"; (Takeuchi et al., *Agents Actions* 36:312-316, (1992)); and metalloproteinase inhibitors such as BB94.

[0949] Additional anti-angiogenic factors that may also be utilized within the context of the present invention include Thalidomide, (Celgene, Warren, NJ); Angiostatic steroid; AGM-1470 (H. Brem and J. Folkman *J Pediatr. Surg.* 28:445-51 (1993)); an integrin alpha v beta 3 antagonist (C. Storgard et al., *J Clin. Invest.* 103:47-54 (1999)); carboxynaminolimidazole; Carboxyamidotriazole (CAI) (National Cancer Institute, Bethesda, MD); Conbretastatin A-4 (CA4P) (OXiGENE, Boston, MA); Squalamine (Magainin Pharmaceuticals, Plymouth Meeting, PA); TNP-470, (Tap Pharmaceuticals, Deerfield, IL); ZD-0101 AstraZeneca (London, UK); APRA (CT2584); Benefin, Byrostatin-1 (SC339555); CGP-41251 (PKC 412); CM101; Dexrazoxane (ICRF187); DMXAA; Endostatin; Flavopridiol; Genestein; GTE; ImmTher; Iressa (ZD1839); Octreotide (Somatostatin); Panretin; Penacillamine; Photopoint; PI-88; Prinomastat (AG-3340) Purlitin; Suradista, (FCE26644); Tamoxifen (Nolvadex); Tazarotene; Tetrathiomolybdate; Xeloda (Capecitabine); and 5-Fluorouracil.

[0950] Anti-angiogenic agents that may be administered in combination with the compounds of the invention may work through a variety of mechanisms including, but not limited to, inhibiting proteolysis of the extracellular matrix, blocking the function of endothelial cell-extracellular matrix adhesion molecules, by antagonizing the function of angiogenesis inducers such as growth factors, and inhibiting integrin receptors expressed on proliferating endothelial cells. Examples of anti-angiogenic inhibitors that interfere with extracellular matrix proteolysis and which may be administered in combination with the compositions of the invention include, but are not limited to, AG-3340 (Agouron, La Jolla, CA), BAY-12-9566 (Bayer, West Haven, CT), BMS-275291 (Bristol Myers Squibb, Princeton, NJ), CGS-27032A (Novartis, East Hanover, NJ), Marimastat (British Biotech, Oxford, UK), and Metastat (Aeterna, St-Foy, Quebec). Examples of anti-angiogenic inhibitors that act by blocking the function of endothelial cell-extracellular matrix adhesion molecules and which may be administered in combination with the compositions of the invention include, but are not limited to, EMD-121974 (Merck KgaA Darmstadt, Germany) and Vitaxin (Ixsys, La Jolla, CA/Medimmune, Gaithersburg, MD). Examples of anti-angiogenic agents that act by directly antagonizing or inhibiting angiogenesis inducers and which may be administered in combination with the compositions of the invention include, but are not limited to, Angiozyme (Ribozyme, Boulder, CO), Anti-VEGF antibody (Genentech, S. San Francisco, CA), PTK-787/ZK-225846 (Novartis, Basel, Switzerland), SU-101 (Sugen, S. San Francisco, CA), SU-5416 (Sugen/ Pharmacia Upjohn, Bridgewater, NJ), and SU-6668 (Sugen). Other anti-angiogenic agents act to indirectly inhibit angiogenesis. Examples of indirect inhibitors of angiogenesis which may be administered in combination with the compositions of the invention include, but are not limited to, IM-862 (Cytran, Kirkland, WA), Interferon-alpha, IL-12 (Roche, Nutley, NJ), and Pentosan polysulfate (Georgetown University, Washington, DC).

[0951] In particular embodiments, the use of compositions of the invention in combination with anti-angiogenic agents is contemplated for the treatment, prevention, and/or amelioration of an autoimmune disease, such as for example, an autoimmune disease described herein.

[0952] In a particular embodiment, the use of compositions of the invention in combination with anti-angiogenic agents is contemplated for the treatment, prevention, and/or amelioration of arthritis. In a more particular embodiment, the use of compositions

of the invention in combination with anti-angiogenic agents is contemplated for the treatment, prevention, and/or amelioration of rheumatoid arthritis.

[0953] In another embodiment, the polynucleotides encoding a polypeptide of the present invention are administered in combination with an angiogenic protein, or polynucleotides encoding an angiogenic protein. Examples of angiogenic proteins that may be administered with the compositions of the invention include, but are not limited to, acidic and basic fibroblast growth factors, VEGF-1, VEGF-2, VEGF-3, epidermal growth factor alpha and beta, platelet-derived endothelial cell growth factor, platelet-derived growth factor, tumor necrosis factor alpha, hepatocyte growth factor, insulin-like growth factor, colony stimulating factor, macrophage colony stimulating factor, granulocyte/macrophage colony stimulating factor, and nitric oxide synthase.

[0954] In additional embodiments, compositions of the invention are administered in combination with a chemotherapeutic agent. Chemotherapeutic agents that may be administered with the Therapeutics of the invention include, but are not limited to alkylating agents such as nitrogen mustards (for example, Mechlorethamine, cyclophosphamide, Cyclophosphamide Ifosfamide, Melphalan (L-sarcosine), and Chlorambucil), ethylenimines and methylmelamines (for example, Hexamethylmelamine and Thiotepa), alkyl sulfonates (for example, Busulfan), nitrosoureas (for example, Carmustine (BCNU), Lomustine (CCNU), Semustine (methyl-CCNU), and Streptozocin (streptozotocin)), triazines (for example, Dacarbazine (DTIC; dimethyltriazenoimidazolecarboxamide)), folic acid analogs (for example, Methotrexate (amethopterin)), pyrimidine analogs (for example, Fluorouracil (5-fluorouracil; 5-FU), Floxuridine (fluorodeoxyuridine; FudR), and Cytarabine (cytosine arabinoside)), purine analogs and related inhibitors (for example, Mercaptopurine (6-mercaptopurine; 6-MP), Thioguanine (6-thioguanine; TG), and Pentostatin (2'-deoxycoformycin)), vinca alkaloids (for example, Vinblastine (VLB, vinblastine sulfate)) and Vincristine (vincristine sulfate)), epipodophyllotoxins (for example, Etoposide and Teniposide), antibiotics (for example, Dactinomycin (actinomycin D), Daunorubicin (daunomycin; rubidomycin), Doxorubicin, Bleomycin, Plicamycin (mithramycin), and Mitomycin (mitomycin C)), enzymes (for example, L-Asparaginase), biological response modifiers (for example, Interferon-alpha and interferon-alpha-2b), platinum coordination compounds (for example, Cisplatin (cis-DDP) and Carboplatin), anthracenedione (Mitoxantrone),

substituted ureas (for example, Hydroxyurea), methylhydrazine derivatives (for example, Procarbazine (N-methylhydrazine; MIH), adrenocorticosteroids (for example, Prednisone), progestins (for example, Hydroxyprogesterone caproate, Medroxyprogesterone, Medroxyprogesterone acetate, and Megestrol acetate), estrogens (for example, Diethylstilbestrol (DES), Diethylstilbestrol diphosphate, Estradiol, and Ethinyl estradiol), antiestrogens (for example, Tamoxifen), androgens (Testosterone propionate, and Fluoxymesterone), antiandrogens (for example, Flutamide), gonadotropin-releasing hormone analogs (for example, Leuprolide), other hormones and hormone analogs (for example, methyltestosterone, estramustine, estramustine phosphate sodium, chlorotrianisene, and testolactone), and others (for example, dicarbazine, glutamic acid, and mitotane).

[0955] In one embodiment, the compositions of the invention are administered in combination with one or more of the following drugs: infliximab (also known as Remicade™ Centocor, Inc.), Trocade (Roche, RO-32-3555), Leflunomide (also known as Arava™ from Hoechst Marion Roussel), Kineret™ (an IL-1 Receptor antagonist also known as Anakinra from Amgen, Inc.)

[0956] In a specific embodiment, compositions of the invention are administered in combination with CHOP (cyclophosphamide, doxorubicin, vincristine, and prednisone) or combination of one or more of the components of CHOP. In one embodiment, the compositions of the invention are administered in combination with anti-CD20 antibodies, human monoclonal anti-CD20 antibodies. In another embodiment, the compositions of the invention are administered in combination with anti-CD20 antibodies and CHOP, or anti-CD20 antibodies and any combination of one or more of the components of CHOP, particularly cyclophosphamide and/or prednisone. In a specific embodiment, compositions of the invention are administered in combination with Rituximab. In a further embodiment, compositions of the invention are administered with Rituximab and CHOP, or Rituximab and any combination of one or more of the components of CHOP, particularly cyclophosphamide and/or prednisone. In a specific embodiment, compositions of the invention are administered in combination with tositumomab. In a further embodiment, compositions of the invention are administered with tositumomab and CHOP, or tositumomab and any combination of one or more of the components of CHOP, particularly cyclophosphamide and/or prednisone. The anti-CD20 antibodies may

optionally be associated with radioisotopes, toxins or cytotoxic prodrugs.

[0957] In another specific embodiment, the compositions of the invention are administered in combination Zevalin™. In a further embodiment, compositions of the invention are administered with Zevalin™ and CHOP, or Zevalin™ and any combination of one or more of the components of CHOP, particularly cyclophosphamide and/or prednisone. Zevalin™ may be associated with one or more radisotopes. Particularly preferred isotopes are ⁹⁰Y and ¹¹¹In.

[0958] In an additional embodiment, the Therapeutics of the invention are administered in combination with cytokines. Cytokines that may be administered with the Therapeutics of the invention include, but are not limited to, IL2, IL3, IL4, IL5, IL6, IL7, IL10, IL12, IL13, IL15, anti-CD40, CD40L, IFN-gamma and TNF-alpha. In another embodiment, Therapeutics of the invention may be administered with any interleukin, including, but not limited to, IL-1alpha, IL-1beta, IL-2, IL-3, IL-4, IL-5, IL-6, IL-7, IL-8, IL-9, IL-10, IL-11, IL-12, IL-13, IL-14, IL-15, IL-16, IL-17, IL-18, IL-19, IL-20, and IL-21.

[0959] In one embodiment, the Therapeutics of the invention are administered in combination with members of the TNF family. TNF, TNF-related or TNF-like molecules that may be administered with the Therapeutics of the invention include, but are not limited to, soluble forms of TNF-alpha, lymphotoxin-alpha (LT-alpha, also known as TNF-beta), LT-beta (found in complex heterotrimer LT-alpha2-beta), OPGL, FasL, CD27L, CD30L, CD40L, 4-1BBL, DcR3, OX40L, TNF-gamma (International Publication No. WO 96/14328), AIM-I (International Publication No. WO 97/33899), endokine-alpha (International Publication No. WO 98/07880), OPG, and neutrokin-alpha (International Publication No. WO 98/18921, OX40, and nerve growth factor (NGF), and soluble forms of Fas, CD30, CD27, CD40 and 4-1BB, TR2 (International Publication No. WO 96/34095), DR3 (International Publication No. WO 97/33904), DR4 (International Publication No. WO 98/32856), TR5 (International Publication No. WO 98/30693), TRANK, TR9 (International Publication No. WO 98/56892), TR10 (International Publication No. WO 98/54202), 312C2 (International Publication No. WO 98/06842), and TR12, and soluble forms CD154, CD70, and CD153.

[0960] In an additional embodiment, the Therapeutics of the invention are administered in combination with angiogenic proteins. Angiogenic proteins that may be administered

with the Therapeutics of the invention include, but are not limited to, Glioma Derived Growth Factor (GDGF), as disclosed in European Patent Number EP-399816; Platelet Derived Growth Factor-A (PDGF-A), as disclosed in European Patent Number EP-682110; Platelet Derived Growth Factor-B (PDGF-B), as disclosed in European Patent Number EP-282317; Placental Growth Factor (PIGF), as disclosed in International Publication Number WO 92/06194; Placental Growth Factor-2 (PIGF-2), as disclosed in Hauser et al., Growth Factors, 4:259-268 (1993); Vascular Endothelial Growth Factor (VEGF), as disclosed in International Publication Number WO 90/13649; Vascular Endothelial Growth Factor-A (VEGF-A), as disclosed in European Patent Number EP-506477; Vascular Endothelial Growth Factor-2 (VEGF-2), as disclosed in International Publication Number WO 96/39515; Vascular Endothelial Growth Factor B (VEGF-3); Vascular Endothelial Growth Factor B-186 (VEGF-B186), as disclosed in International Publication Number WO 96/26736; Vascular Endothelial Growth Factor-D (VEGF-D), as disclosed in International Publication Number WO 98/02543; Vascular Endothelial Growth Factor-D (VEGF-D), as disclosed in International Publication Number WO 98/07832; and Vascular Endothelial Growth Factor-E (VEGF-E), as disclosed in German Patent Number DE19639601. The above mentioned references are herein incorporated by reference in their entireties.

[0961] In an additional embodiment, the Therapeutics of the invention are administered in combination with Fibroblast Growth Factors. Fibroblast Growth Factors that may be administered with the Therapeutics of the invention include, but are not limited to, FGF-1, FGF-2, FGF-3, FGF-4, FGF-5, FGF-6, FGF-7, FGF-8, FGF-9, FGF-10, FGF-11, FGF-12, FGF-13, FGF-14, and FGF-15.

[0962] In an additional embodiment, the Therapeutics of the invention are administered in combination with hematopoietic growth factors. Hematopoietic growth factors that may be administered with the Therapeutics of the invention include, but are not limited to, granulocyte macrophage colony stimulating factor (GM-CSF) (sargramostim, LEUKINE™, PROKINE™), granulocyte colony stimulating factor (G-CSF) (filgrastim, NEUPOGEN™), macrophage colony stimulating factor (M-CSF, CSF-1) erythropoietin (epoetin alfa, EPOGEN™, PROCRIT™), stem cell factor (SCF, c-kit ligand, steel factor), megakaryocyte colony stimulating factor, PIXY321 (a GMCSF/IL-3

fusion protein), interleukins, especially any one or more of IL-1 through IL-12, interferon-gamma, or thrombopoietin.

[0963] In certain embodiments, Therapeutics of the present invention are administered in combination with adrenergic blockers, such as, for example, acebutolol, atenolol, betaxolol, bisoprolol, carteolol, labetalol, metoprolol, nadolol, oxprenolol, penbutolol, pindolol, propranolol, sotalol, and timolol.

[0964] In another embodiment, the Therapeutics of the invention are administered in combination with an antiarrhythmic drug (e.g., adenosine, amiodarone, bretylium, digitalis, digoxin, digitoxin, diltiazem, disopyramide, esmolol, flecainide, lidocaine, mexiletine, moricizine, phenytoin, procainamide, N-acetyl procainamide, propafenone, propranolol, quinidine, sotalol, tocainide, and verapamil).

[0965] In another embodiment, the Therapeutics of the invention are administered in combination with diuretic agents, such as carbonic anhydrase-inhibiting agents (e.g., acetazolamide, dichlorphenamide, and methazolamide), osmotic diuretics (e.g., glycerin, isosorbide, mannitol, and urea), diuretics that inhibit $\text{Na}^+\text{-K}^+\text{-2Cl}^-$ symport (e.g., furosemide, bumetanide, azosemide, piretanide, tripamide, ethacrynic acid, muzolimine, and torsemide), thiazide and thiazide-like diuretics (e.g., bendroflumethiazide, benzthiazide, chlorothiazide, hydrochlorothiazide, hydroflumethiazide, methyclothiazide, polythiazide, trichormethiazide, chlorthalidone, indapamide, metolazone, and quinethazone), potassium sparing diuretics (e.g., amiloride and triamterene); and mineralocorticoid receptor antagonists (e.g., spironolactone, canrenone, and potassium canrenoate).

[0966] In one embodiment, the Therapeutics of the invention are administered in combination with treatments for endocrine and/or hormone imbalance disorders. Treatments for endocrine and/or hormone imbalance disorders include, but are not limited to, ^{127}I , radioactive isotopes of iodine such as ^{131}I and ^{123}I ; recombinant growth hormone, such as HUMATROPE™ (recombinant somatropin); growth hormone analogs such as PROTROPIN™ (somatrem); dopamine agonists such as PARLODEL™ (bromocriptine); somatostatin analogs such as SANDOSTATIN™ (octreotide); gonadotropin preparations such as PREGNYL™, A.P.L.™ and PROFASI™ (chorionic gonadotropin (CG)), PERGONAL™ (menotropins), and METRODIN™ (urofollitropin (uFSH)); synthetic human gonadotropin releasing hormone preparations such as FACTREL™ and

LUTREPULSE™ (gonadorelin hydrochloride); synthetic gonadotropin agonists such as LUPRON™ (leuprolide acetate), SUPPRELIN™ (histrelin acetate), SYNAREL™ (nafarelin acetate), and ZOLADEX™ (goserelin acetate); synthetic preparations of thyrotropin-releasing hormone such as RELEFACT TRH™ and THYPINONE™ (protirelin); recombinant human TSH such as THYROGEN™; synthetic preparations of the sodium salts of the natural isomers of thyroid hormones such as L-T₄™, SYNTHROID™ and LEVOTHROID™ (levothyroxine sodium), L-T₃™, CYTOMEL™ and TRIOSTAT™ (liothyroine sodium), and THYROLAR™ (liotrix); antithyroid compounds such as 6-*n*-propylthiouracil (propylthiouracil), 1-methyl-2-mercaptoimidazole and TAPAZOLE™ (methimazole), NEO-MERCAZOLE™ (carbimazole); beta-adrenergic receptor antagonists such as propranolol and esmolol; Ca²⁺ channel blockers; dexamethasone and iodinated radiological contrast agents such as TELEPAQUE™ (iopanoic acid) and ORAGRAFIN™ (sodium ipodate).

[0967] Additional treatments for endocrine and/or hormone imbalance disorders include, but are not limited to, estrogens or conjugated estrogens such as ESTRACE™ (estradiol), ESTINYL™ (ethinyl estradiol), PREMARIN™, ESTRATAB™, ORTHO-EST™, OGEN™ and estropipate (estrone), ESTROVIS™ (quinestrol), ESTRADERM™ (estradiol), DELESTROGEN™ and VALERGEN™ (estradiol valerate), DEPO-ESTRADIOL CYPIONATE™ and ESTROJECT LA™ (estradiol cypionate); antiestrogens such as NOLVADEX™ (tamoxifen), SEROPHENE™ and CLOMID™ (clomiphene); progestins such as DURALUTIN™ (hydroxyprogesterone caproate), MPA™ and DEPO-PROVERA™ (medroxyprogesterone acetate), PROVERA™ and CYCRIN™ (MPA), MEGACE™ (megestrol acetate), NORLUTIN™ (norethindrone), and NORLUTATE™ and AYGESTIN™ (norethindrone acetate); progesterone implants such as NORPLANT SYSTEM™ (subdermal implants of norgestrel); antiprogestins such as RU 486™ (mifepristone); hormonal contraceptives such as ENOVID™ (norethynodrel plus mestranol), PROGESTASERT™ (intrauterine device that releases progesterone), LOESTRIN™, BREVICON™, MODICON™, GENORA™, NELONA™, NORINYL™, OVACON-35™ and OVACON-50™ (ethinyl estradiol/norethindrone), LEVLEN™, NORDETTE™, TRI-LEVLEN™ and TRIPHASIL-21™ (ethinyl estradiol/levonorgestrel)

LO/OVRAL™ and OVRAL™ (ethinyl estradiol/norgestrel), DEMULEN™ (ethinyl estradiol/ethynodiol diacetate), NORINYL™, ORTHO-NOVUM™, NORETHIN™, GENORA™, and NELOVA™ (norethindrone/mestranol), DESOGEN™ and ORTHO-CEPT™ (ethinyl estradiol/desogestrel), ORTHO-CYCLEN™ and ORTHO-TRICYCLEN™ (ethinyl estradiol/norgestimate), MICRONOR™ and NOR-QD™ (norethindrone), and OVRETTE™ (norgestrel).

[0968] Additional treatments for endocrine and/or hormone imbalance disorders include, but are not limited to, testosterone esters such as methenolone acetate and testosterone undecanoate; parenteral and oral androgens such as TESTOJECT-50™ (testosterone), TESTEX™ (testosterone propionate), DELATESTRYL™ (testosterone enanthate), DEPO-TESTOSTERONE™ (testosterone cypionate), DANOCRINE™ (danazol), HALOTESTIN™ (fluoxymesterone), ORETON METHYL™, TESTRED™ and VIRILON™ (methyltestosterone), and OXANDRIN™ (oxandrolone); testosterone transdermal systems such as TESTODERM™; androgen receptor antagonist and 5-alpha-reductase inhibitors such as ANDROCUR™ (cyproterone acetate), EULEXIN™ (flutamide), and PROSCAR™ (finasteride); adrenocorticotrophic hormone preparations such as CORTROSYN™ (cosyntropin); adrenocortical steroids and their synthetic analogs such as ACLOVATE™ (alclometasone dipropionate), CYCLOCORT™ (amcinonide), BECLOVENT™ and VANCERIL™ (beclomethasone dipropionate), CELESTONE™ (betamethasone), BENISONE™ and UTICORT™ (betamethasone benzoate), DIPROSONE™ (betamethasone dipropionate), CELESTONE PHOSPHATE™ (betamethasone sodium phosphate), CELESTONE SOLUSPAN™ (betamethasone sodium phosphate and acetate), BETA-VAL™ and VALISONE™ (betamethasone valerate), TEMOVATE™ (clobetasol propionate), CLODERM™ (clocortolone pivalate), CORTEF™ and HYDROCORTONE™ (cortisol (hydrocortisone)), HYDROCORTONE ACETATE™ (cortisol (hydrocortisone) acetate), LOCOID™ (cortisol (hydrocortisone) butyrate), HYDROCORTONE PHOSPHATE™ (cortisol (hydrocortisone) sodium phosphate), A-HYDROCORT™ and SOLU CORTEF™ (cortisol (hydrocortisone) sodium succinate), WESTCORT™ (cortisol (hydrocortisone) valerate), CORTISONE ACETATE™ (cortisone acetate), DESOWEN™ and TRIDESILON™ (desonide),

TOPICORT™ (desoximetasone), DECADRON™ (dexamethasone), DECADRON LA™ (dexamethasone acetate), DECADRON PHOSPHATE™ and HEXADROL PHOSPHATE™ (dexamethasone sodium phosphate), FLORONE™ and MAXIFLOR™ (diflorasone diacetate), FLORINEF ACETATE™ (fludrocortisone acetate), AEROBID™ and NASALIDE™ (flunisolide), FLUONID™ and SYNALAR™ (fluocinolone acetonide), LIDEX™ (fluocinonide), FLUOR-OP™ and FML™ (fluorometholone), CORDRAN™ (flurandrenolide), HALOG™ (halcinonide), HMS LIZUIFILM™ (medrysone), MEDROL™ (methylprednisolone), DEPO-MEDROL™ and MEDROL ACETATE™ (methylprednisone acetate), A-METHAPRED™ and SOLUMEDROL™ (methylprednisolone sodium succinate), ELOCON™ (mometasone furoate), HALDRONE™ (paramethasone acetate), DELTA-CORTEF™ (prednisolone), ECONOPRED™ (prednisolone acetate), HYDELTRASOL™ (prednisolone sodium phosphate), HYDELTRA-T.B.A™ (prednisolone tebutate), DELTASONE™ (prednisone), ARISTOCORT™ and KENACORT™ (triamcinolone), KENALOG™ (triamcinolone acetonide), ARISTOCORT™ and KENACORT DIACETATE™ (triamcinolone diacetate), and ARISTOSPAN™ (triamcinolone hexacetonide); inhibitors of biosynthesis and action of adrenocortical steroids such as CYTADREN™ (aminoglutethimide), NIZORAL™ (ketoconazole), MODRASTANE™ (trilostane), and METOPIRONE™ (metyrapone); bovine, porcine or human insulin or mixtures thereof; insulin analogs; recombinant human insulin such as HUMULIN™ and NOVOLIN™; oral hypoglycemic agents such as ORAMIDE™ and ORINASE™ (tolbutamide), DIABINESE™ (chlorpropamide), TOLAMIDE™ and TOLINASE™ (tolazamide), DYMELOR™ (acetohexamide), glibenclamide, MICRONASE™, DIBETA™ and GLYNASE™ (glyburide), GLUCOTROL™ (glipizide), and DIAMICRON™ (gliclazide), GLUCOPHAGE™ (metformin), ciglitazone, pioglitazone, and alpha-glucosidase inhibitors; bovine or porcine glucagon; somatostatins such as SANDOSTATIN™ (octreotide); and diazoxides such as PROGLYCEM™ (diazoxide).

[0969] In one embodiment, the Therapeutics of the invention are administered in combination with treatments for uterine motility disorders. Treatments for uterine motility disorders include, but are not limited to, estrogen drugs such as conjugated estrogens (e.g., PREMARIN® and ESTRATAB®), estradiols (e.g., CLIMARA® and ALORA®),

estropipate, and chlorotrianisene; progestin drugs (e.g., AMEN[®] (medroxyprogesterone), MICRONOR[®] (norethidrone acetate), PROMETRIUM[®] progesterone, and megestrol acetate); and estrogen/progesterone combination therapies such as, for example, conjugated estrogens/medroxyprogesterone (e.g., PREMPRO[™] and PREMPHASE[®]) and norethindrone acetate/ethinyl estsradiol (e.g., FEMHRT[™]).

[0970] In an additional embodiment, the Therapeutics of the invention are administered in combination with drugs effective in treating iron deficiency and hypochromic anemias, including but not limited to, ferrous sulfate (iron sulfate, FEOSOL[™]), ferrous fumarate (e.g., FEOSTAT[™]), ferrous gluconate (e.g., FERGON[™]), polysaccharide-iron complex (e.g., NIFEREX[™]), iron dextran injection (e.g., INFED[™]), cupric sulfate, pyroxidine, riboflavin, Vitamin B₁₂, cyancobalamin injection (e.g., REDISOL[™], RUBRAMIN PCT[™]), hydroxocobalamin, folic acid (e.g., FOLVITE[™]), leucovorin (folinic acid, 5-CHOH4PteGlu, citrovorum factor) or WELLCOVORIN (Calcium salt of leucovorin), transferrin or ferritin.

[0971] In certain embodiments, the Therapeutics of the invention are administered in combination with agents used to treat psychiatric disorders. Psychiatric drugs that may be administered with the Therapeutics of the invention include, but are not limited to, antipsychotic agents (e.g., chlorpromazine, chlorprothixene, clozapine, fluphenazine, haloperidol, loxapine, mesoridazine, molindone, olanzapine, perphenazine, pimozide, quetiapine, risperidone, thioridazine, thiothixene, trifluoperazine, and triflupromazine), antimanic agents (e.g., carbamazepine, divalproex sodium, lithium carbonate, and lithium citrate), antidepressants (e.g., amitriptyline, amoxapine, bupropion, citalopram, clomipramine, desipramine, doxepin, fluvoxamine, fluoxetine, imipramine, isocarboxazid, maprotiline, mirtazapine, nefazodone, nortriptyline, paroxetine, phenelzine, protriptyline, sertraline, tranylcypromine, trazodone, trimipramine, and venlafaxine), antianxiety agents (e.g., alprazolam, buspirone, chlordiazepoxide, clorazepate, diazepam, halazepam, lorazepam, oxazepam, and prazepam), and stimulants (e.g., d-amphetamine, methylphenidate, and pemoline).

[0972] In other embodiments, the Therapeutics of the invention are administered in combination with agents used to treat neurological disorders. Neurological agents that may be administered with the Therapeutics of the invention include, but are not limited to, antiepileptic agents (e.g., carbamazepine, clonazepam, ethosuximide, phenobarbital,

phenytoin, primidone, valproic acid, divalproex sodium, felbamate, gabapentin, lamotrigine, levetiracetam, oxcarbazepine, tiagabine, topiramate, zonisamide, diazepam, lorazepam, and clonazepam), antiparkinsonian agents (e.g., levodopa/carbidopa, selegiline, amantidine, bromocriptine, pergolide, ropinirole, pramipexole, benzotropine; biperiden; ethopropazine; procyclidine; trihexyphenidyl, tolcapone), and ALS therapeutics (e.g. riluzole).

[0973] In another embodiment, Therapeutics of the invention are administered in combination with vasodilating agents and/or calcium channel blocking agents. Vasodilating agents that may be administered with the Therapeutics of the invention include, but are not limited to, Angiotensin Converting Enzyme (ACE) inhibitors (e.g., papaverine, isoxsuprine, benazepril, captopril, cilazapril, enalapril, enalaprilat, fosinopril, lisinopril, moexipril, perindopril, quinapril, ramipril, spirapril, trandolapril, and nylidrin), and nitrates (e.g., isosorbide dinitrate, isosorbide mononitrate, and nitroglycerin). Examples of calcium channel blocking agents that may be administered in combination with the Therapeutics of the invention include, but are not limited to amlodipine, bepridil, diltiazem, felodipine, flunarizine, isradipine, nicardipine, nifedipine, nimodipine, and verapamil.

[0974] In certain embodiments, the Therapeutics of the invention are administered in combination with treatments for gastrointestinal disorders. Treatments for gastrointestinal disorders that may be administered with the Therapeutic of the invention include, but are not limited to, H₂ histamine receptor antagonists (e.g., TAGAMETTM (cimetidine), ZANTACTM (ranitidine), PEPCIDTM (famotidine), and AXIDTM (nizatidine)); inhibitors of H⁺, K⁺ ATPase (e.g., PREVACIDTM (lansoprazole) and PRILOSECTM (omeprazole)); Bismuth compounds (e.g., PEPTO-BISMOLTM (bismuth subsalicylate) and DE-NOLTM (bismuth subcitrate)); various antacids; sucralfate; prostaglandin analogs (e.g. CYTOTECTM (misoprostol)); muscarinic cholinergic antagonists; laxatives (e.g., surfactant laxatives, stimulant laxatives, saline and osmotic laxatives); antidiarrheal agents (e.g., LOMOTILTM (diphenoxylate), MOTOFENTM (diphenoxin), and IMODIUMTM (loperamide hydrochloride)), synthetic analogs of somatostatin such as SANDOSTATINTM (octreotide), antiemetic agents (e.g., ZOFRANTM (ondansetron), KYTRILTM (granisetron hydrochloride), tropisetron, dolasetron, metoclopramide, chlorpromazine, perphenazine, prochlorperazine, promethazine, thiethylperazine,

triflupromazine, domperidone, haloperidol, droperidol, trimethobenzamide, dexamethasone, methylprednisolone, dronabinol, and nabilone); D2 antagonists (e.g., metoclopramide, trimethobenzamide and chlorpromazine); bile salts; chenodeoxycholic acid; ursodeoxycholic acid; and pancreatic enzyme preparations such as pancreatin and pancrelipase.

[0975] In additional embodiments, the Therapeutics of the invention are administered in combination with other therapeutic or prophylactic regimens, such as, for example, radiation therapy.

Example 14: Method of Treating Decreased Levels of the Polypeptide

[0976] The present invention relates to a method for treating an individual in need of an increased level of a polypeptide of the invention in the body comprising administering to such an individual a composition comprising a therapeutically effective amount of an agonist of the invention (including polypeptides of the invention). Moreover, it will be appreciated that conditions caused by a decrease in the standard or normal expression level of a polypeptide of the present invention in an individual can be treated by administering the agonist or antagonist of the present invention. Thus, the invention also provides a method of treatment of an individual in need of an increased level of the polypeptide comprising administering to such an individual a Therapeutic comprising an amount of the agonist or antagonist to increase the activity level of the polypeptide in such an individual.

[0977] For example, a patient with decreased levels of a polypeptide receives a daily dose 0.1-100 ug/kg of the agonist or antagonist for six consecutive days. The exact details of the dosing scheme, based on administration and formulation, are provided in Example 13.

Example 15: Method of Treating Increased Levels of the Polypeptide

[0978] The present invention also relates to a method of treating an individual in need of a decreased level of a polypeptide of the invention in the body comprising

administering to such an individual a composition comprising a therapeutically effective amount of an antagonist of the invention (including polypeptides and antibodies of the invention).

[0979] In one example, antisense technology is used to inhibit production of a polypeptide of the present invention. This technology is one example of a method of decreasing levels of a polypeptide, due to a variety of etiologies, such as cancer.

[0980] For example, a patient diagnosed with abnormally increased levels of a polypeptide is administered intravenously antisense polynucleotides at 0.5, 1.0, 1.5, 2.0 and 3.0 mg/kg day for 21 days. This treatment is repeated after a 7-day rest period if the treatment was well tolerated. The antisense polynucleotides of the present invention can be formulated using techniques and formulations described herein (e.g. see Example 13), or otherwise known in the art.

Example 16: Method of Treatment Using Gene Therapy-Ex Vivo

[0981] One method of gene therapy transplants fibroblasts, which are capable of expressing a polypeptide, onto a patient. Generally, fibroblasts are obtained from a subject by skin biopsy. The resulting tissue is placed in tissue-culture medium and separated into small pieces. Small chunks of the tissue are placed on a wet surface of a tissue culture flask, approximately ten pieces are placed in each flask. The flask is turned upside down, closed tight and left at room temperature over night. After 24 hours at room temperature, the flask is inverted and the chunks of tissue remain fixed to the bottom of the flask and fresh media (e.g., Ham's F12 media, with 10% FBS, penicillin and streptomycin) is added. The flasks are then incubated at 37 degree C for approximately one week.

[0982] At this time, fresh media is added and subsequently changed every several days. After an additional two weeks in culture, a monolayer of fibroblasts emerge. The monolayer is trypsinized and scaled into larger flasks.

[0983] pMV-7 (Kirschmeier, P.T. et al., DNA, 7:219-25 (1988)), flanked by the long terminal repeats of the Moloney murine sarcoma virus, is digested with EcoRI and HindIII and subsequently treated with calf intestinal phosphatase. The linear vector is fractionated on agarose gel and purified, using glass beads.

[0984] The cDNA encoding a polypeptide of the present invention can be amplified using PCR primers which correspond to the 5' and 3' end sequences respectively as set forth in Example 1 using primers and having appropriate restriction sites and initiation/stop codons, if necessary. Preferably, the 5' primer contains an EcoRI site and the 3' primer includes a HindIII site. Equal quantities of the Moloney murine sarcoma virus linear backbone and the amplified EcoRI and HindIII fragment are added together, in the presence of T4 DNA ligase. The resulting mixture is maintained under conditions appropriate for ligation of the two fragments. The ligation mixture is then used to transform bacteria HB101, which are then plated onto agar containing kanamycin for the purpose of confirming that the vector has the gene of interest properly inserted.

[0985] The amphotropic pA317 or GP+am12 packaging cells are grown in tissue culture to confluent density in Dulbecco's Modified Eagles Medium (DMEM) with 10% calf serum (CS), penicillin and streptomycin. The MSV vector containing the gene is then added to the media and the packaging cells transduced with the vector. The packaging cells now produce infectious viral particles containing the gene (the packaging cells are now referred to as producer cells).

[0986] Fresh media is added to the transduced producer cells, and subsequently, the media is harvested from a 10 cm plate of confluent producer cells. The spent media, containing the infectious viral particles, is filtered through a millipore filter to remove detached producer cells and this media is then used to infect fibroblast cells. Media is removed from a sub-confluent plate of fibroblasts and quickly replaced with the media from the producer cells. This media is removed and replaced with fresh media. If the titer of virus is high, then virtually all fibroblasts will be infected and no selection is required. If the titer is very low, then it is necessary to use a retroviral vector that has a selectable marker, such as neo or his. Once the fibroblasts have been efficiently infected, the fibroblasts are analyzed to determine whether protein is produced.

[0987] The engineered fibroblasts are then transplanted onto the host, either alone or after having been grown to confluence on cytodex 3 microcarrier beads.

***Example 17: Gene Therapy Using Endogenous Genes Corresponding To
Polynucleotides of the Invention***

[0988] Another method of gene therapy according to the present invention involves operably associating the endogenous polynucleotide sequence of the invention with a promoter via homologous recombination as described, for example, in U.S. Patent NO: 5,641,670, issued June 24, 1997; International Publication NO: WO 96/29411, published September 26, 1996; International Publication NO: WO 94/12650, published August 4, 1994; Koller et al., *Proc. Natl. Acad. Sci. USA*, 86:8932-8935 (1989); and Zijlstra et al., *Nature*, 342:435-438 (1989). This method involves the activation of a gene which is present in the target cells, but which is not expressed in the cells, or is expressed at a lower level than desired.

[0989] Polynucleotide constructs are made which contain a promoter and targeting sequences, which are homologous to the 5' non-coding sequence of endogenous polynucleotide sequence, flanking the promoter. The targeting sequence will be sufficiently near the 5' end of the polynucleotide sequence so the promoter will be operably linked to the endogenous sequence upon homologous recombination. The promoter and the targeting sequences can be amplified using PCR. Preferably, the amplified promoter contains distinct restriction enzyme sites on the 5' and 3' ends. Preferably, the 3' end of the first targeting sequence contains the same restriction enzyme site as the 5' end of the amplified promoter and the 5' end of the second targeting sequence contains the same restriction site as the 3' end of the amplified promoter.

[0990] The amplified promoter and the amplified targeting sequences are digested with the appropriate restriction enzymes and subsequently treated with calf intestinal phosphatase. The digested promoter and digested targeting sequences are added together in the presence of T4 DNA ligase. The resulting mixture is maintained under conditions appropriate for ligation of the two fragments. The construct is size fractionated on an agarose gel, then purified by phenol extraction and ethanol precipitation.

[0991] In this Example, the polynucleotide constructs are administered as naked polynucleotides via electroporation. However, the polynucleotide constructs may also be administered with transfection-facilitating agents, such as liposomes, viral sequences, viral particles, precipitating agents, etc. Such methods of delivery are known in the art.

[0992] Once the cells are transfected, homologous recombination will take place which results in the promoter being operably linked to the endogenous polynucleotide sequence. This results in the expression of polynucleotide corresponding to the polynucleotide in the cell. Expression may be detected by immunological staining, or any other method known in the art.

[0993] Fibroblasts are obtained from a subject by skin biopsy. The resulting tissue is placed in DMEM + 10% fetal calf serum. Exponentially growing or early stationary phase fibroblasts are trypsinized and rinsed from the plastic surface with nutrient medium. An aliquot of the cell suspension is removed for counting, and the remaining cells are subjected to centrifugation. The supernatant is aspirated and the pellet is resuspended in 5 ml of electroporation buffer (20 mM HEPES pH 7.3, 137 mM NaCl, 5 mM KCl, 0.7 mM Na₂ HPO₄, 6 mM dextrose). The cells are recentrifuged, the supernatant aspirated, and the cells resuspended in electroporation buffer containing 1 mg/ml acetylated bovine serum albumin. The final cell suspension contains approximately 3×10^6 cells/ml. Electroporation should be performed immediately following resuspension.

[0994] Plasmid DNA is prepared according to standard techniques. For example, to construct a plasmid for targeting to the locus corresponding to the polynucleotide of the invention, plasmid pUC18 (MBI Fermentas, Amherst, NY) is digested with HindIII. The CMV promoter is amplified by PCR with an XbaI site on the 5' end and a BamHI site on the 3' end. Two non-coding sequences are amplified via PCR: one non-coding sequence (fragment 1) is amplified with a HindIII site at the 5' end and an Xba site at the 3' end; the other non-coding sequence (fragment 2) is amplified with a BamHI site at the 5' end and a HindIII site at the 3' end. The CMV promoter and the fragments (1 and 2) are digested with the appropriate enzymes (CMV promoter - XbaI and BamHI; fragment 1 - XbaI; fragment 2 - BamHI) and ligated together. The resulting ligation product is digested with HindIII, and ligated with the HindIII-digested pUC18 plasmid.

[0995] Plasmid DNA is added to a sterile cuvette with a 0.4 cm electrode gap (Bio-Rad). The final DNA concentration is generally at least 120 μ g/ml. 0.5 ml of the cell suspension (containing approximately 1.5×10^6 cells) is then added to the cuvette, and the cell suspension and DNA solutions are gently mixed. Electroporation is performed with a Gene-Pulser apparatus (Bio-Rad). Capacitance and voltage are set at 960 μ F and 250-300 V, respectively. As voltage increases, cell survival decreases, but the percentage of

surviving cells that stably incorporate the introduced DNA into their genome increases dramatically. Given these parameters, a pulse time of approximately 14-20 mSec should be observed.

[0996] Electroporated cells are maintained at room temperature for approximately 5 min, and the contents of the cuvette are then gently removed with a sterile transfer pipette. The cells are added directly to 10 ml of prewarmed nutrient media (DMEM with 15% calf serum) in a 10 cm dish and incubated at 37 degree C. The following day, the media is aspirated and replaced with 10 ml of fresh media and incubated for a further 16-24 hours.

[0997] The engineered fibroblasts are then injected into the host, either alone or after having been grown to confluence on cytodex 3 microcarrier beads. The fibroblasts now produce the protein product. The fibroblasts can then be introduced into a patient as described above.

Example 18: Method of Treatment Using Gene Therapy - In Vivo

[0998] Another aspect of the present invention is using *in vivo* gene therapy methods to treat disorders, diseases and conditions. The gene therapy method relates to the introduction of naked nucleic acid (DNA, RNA, and antisense DNA or RNA) sequences into an animal to increase or decrease the expression of the polypeptide. The polynucleotide of the present invention may be operatively linked to (i.e., associated with) a promoter or any other genetic elements necessary for the expression of the polypeptide by the target tissue. Such gene therapy and delivery techniques and methods are known in the art, see, for example, WO90/11092, WO98/11779; U.S. Patent NO. 5693622, 5705151, 5580859; Tabata et al., Cardiovasc. Res. 35(3):470-479 (1997); Chao et al., Pharmacol. Res. 35(6):517-522 (1997); Wolff, Neuromuscul. Disord. 7(5):314-318 (1997); Schwartz et al., Gene Ther. 3(5):405-411 (1996); Tsurumi et al., Circulation 94(12):3281-3290 (1996) (incorporated herein by reference).

[0999] The polynucleotide constructs may be delivered by any method that delivers injectable materials to the cells of an animal, such as, injection into the interstitial space of tissues (heart, muscle, skin, lung, liver, intestine and the like). The polynucleotide constructs can be delivered in a pharmaceutically acceptable liquid or aqueous carrier.

[1000] The term "naked" polynucleotide, DNA or RNA, refers to sequences that are

free from any delivery vehicle that acts to assist, promote, or facilitate entry into the cell, including viral sequences, viral particles, liposome formulations, lipofectin or precipitating agents and the like. However, the polynucleotides of the present invention may also be delivered in liposome formulations (such as those taught in Felgner P.L. et al. (1995) Ann. NY Acad. Sci. 772:126-139 and Abdallah B. et al. (1995) Biol. Cell 85(1):1-7) which can be prepared by methods well known to those skilled in the art.

[1001] The polynucleotide vector constructs used in the gene therapy method are preferably constructs that will not integrate into the host genome nor will they contain sequences that allow for replication. Any strong promoter known to those skilled in the art can be used for driving the expression of DNA. Unlike other gene therapy techniques, one major advantage of introducing naked nucleic acid sequences into target cells is the transitory nature of the polynucleotide synthesis in the cells. Studies have shown that non-replicating DNA sequences can be introduced into cells to provide production of the desired polypeptide for periods of up to six months.

[1002] The polynucleotide construct can be delivered to the interstitial space of tissues within an animal, including muscle, skin, brain, lung, liver, spleen, bone marrow, thymus, heart, lymph, blood, bone, cartilage, pancreas, kidney, gall bladder, stomach, intestine, testis, ovary, uterus, rectum, nervous system, eye, gland, and connective tissue. Interstitial space of the tissues comprises the intercellular fluid, mucopolysaccharide matrix among the reticular fibers of organ tissues, elastic fibers in the walls of vessels or chambers, collagen fibers of fibrous tissues, or that same matrix within connective tissue ensheathing muscle cells or in the lacunae of bone. It is similarly the space occupied by the plasma of the circulation and the lymph fluid of the lymphatic channels. Delivery to the interstitial space of muscle tissue is preferred for the reasons discussed below. They may be conveniently delivered by injection into the tissues comprising these cells. They are preferably delivered to and expressed in persistent, non-dividing cells which are differentiated, although delivery and expression may be achieved in non-differentiated or less completely differentiated cells, such as, for example, stem cells of blood or skin fibroblasts. *In vivo* muscle cells are particularly competent in their ability to take up and express polynucleotides.

[1003] For the naked polynucleotide injection, an effective dosage amount of DNA or RNA will be in the range of from about 0.05 g/kg body weight to about 50 mg/kg body

weight. Preferably the dosage will be from about 0.005 mg/kg to about 20 mg/kg and more preferably from about 0.05 mg/kg to about 5 mg/kg. Of course, as the artisan of ordinary skill will appreciate, this dosage will vary according to the tissue site of injection. The appropriate and effective dosage of nucleic acid sequence can readily be determined by those of ordinary skill in the art and may depend on the condition being treated and the route of administration. The preferred route of administration is by the parenteral route of injection into the interstitial space of tissues. However, other parenteral routes may also be used, such as, inhalation of an aerosol formulation particularly for delivery to lungs or bronchial tissues, throat or mucous membranes of the nose. In addition, naked polynucleotide constructs can be delivered to arteries during angioplasty by the catheter used in the procedure.

[1004] The dose response effects of injected polynucleotide in muscle *in vivo* is determined as follows. Suitable template DNA for production of mRNA coding for polypeptide of the present invention is prepared in accordance with a standard recombinant DNA methodology. The template DNA, which may be either circular or linear, is either used as naked DNA or complexed with liposomes. The quadriceps muscles of mice are then injected with various amounts of the template DNA.

[1005] Five to six week old female and male Balb/C mice are anesthetized by intraperitoneal injection with 0.3 ml of 2.5% Avertin. A 1.5 cm incision is made on the anterior thigh, and the quadriceps muscle is directly visualized. The template DNA is injected in 0.1 ml of carrier in a 1 cc syringe through a 27 gauge needle over one minute, approximately 0.5 cm from the distal insertion site of the muscle into the knee and about 0.2 cm deep. A suture is placed over the injection site for future localization, and the skin is closed with stainless steel clips.

[1006] After an appropriate incubation time (e.g., 7 days) muscle extracts are prepared by excising the entire quadriceps. Every fifth 15 μ m cross-section of the individual quadriceps muscles is histochemically stained for protein expression. A time course for protein expression may be done in a similar fashion except that quadriceps from different mice are harvested at different times. Persistence of DNA in muscle following injection may be determined by Southern blot analysis after preparing total cellular DNA and HIRT supernatants from injected and control mice. The results of the above experimentation in mice can be used to extrapolate proper dosages and other treatment parameters in humans

and other animals using naked DNA.

Example 19: Transgenic Animals

[1007] The polypeptides of the invention can also be expressed in transgenic animals. Animals of any species, including, but not limited to, mice, rats, rabbits, hamsters, guinea pigs, pigs, micro-pigs, goats, sheep, cows and non-human primates, *e.g.*, baboons, monkeys, and chimpanzees may be used to generate transgenic animals. In a specific embodiment, techniques described herein or otherwise known in the art, are used to express polypeptides of the invention in humans, as part of a gene therapy protocol.

[1008] Any technique known in the art may be used to introduce the transgene (*i.e.*, polynucleotides of the invention) into animals to produce the founder lines of transgenic animals. Such techniques include, but are not limited to, pronuclear microinjection (Paterson et al., Appl. Microbiol. Biotechnol. 40:691-698 (1994); Carver et al., Biotechnology (NY) 11:1263-1270 (1993); Wright et al., Biotechnology (NY) 9:830-834 (1991); and Hoppe et al., U.S. Pat. No. 4,873,191 (1989)); retrovirus mediated gene transfer into germ lines (Van der Putten et al., Proc. Natl. Acad. Sci., USA 82:6148-6152 (1985)), blastocysts or embryos; gene targeting in embryonic stem cells (Thompson et al., Cell 56:313-321 (1989)); electroporation of cells or embryos (Lo, 1983, Mol Cell. Biol. 3:1803-1814 (1983)); introduction of the polynucleotides of the invention using a gene gun (see, *e.g.*, Ulmer et al., Science 259:1745 (1993); introducing nucleic acid constructs into embryonic pluripotent stem cells and transferring the stem cells back into the blastocyst; and sperm-mediated gene transfer (Lavitrano et al., Cell 57:717-723 (1989); etc. For a review of such techniques, see Gordon, "Transgenic Animals," Intl. Rev. Cytol. 115:171-229 (1989), which is incorporated by reference herein in its entirety.

[1009] Any technique known in the art may be used to produce transgenic clones containing polynucleotides of the invention, for example, nuclear transfer into enucleated oocytes of nuclei from cultured embryonic, fetal, or adult cells induced to quiescence (Campbell et al., Nature 380:64-66 (1996); Wilmut et al., Nature 385:810-813 (1997)).

[1010] The present invention provides for transgenic animals that carry the transgene in all their cells, as well as animals which carry the transgene in some, but not all their cells, *i.e.*, mosaic animals or chimeric. The transgene may be integrated as a single

transgene or as multiple copies such as in concatamers, *e.g.*, head-to-head tandems or head-to-tail tandems. The transgene may also be selectively introduced into and activated in a particular cell type by following, for example, the teaching of Lasko et al. (Lasko et al., Proc. Natl. Acad. Sci. USA 89:6232-6236 (1992)). The regulatory sequences required for such a cell-type specific activation will depend upon the particular cell type of interest, and will be apparent to those of skill in the art. When it is desired that the polynucleotide transgene be integrated into the chromosomal site of the endogenous gene, gene targeting is preferred. Briefly, when such a technique is to be utilized, vectors containing some nucleotide sequences homologous to the endogenous gene are designed for the purpose of integrating, via homologous recombination with chromosomal sequences, into and disrupting the function of the nucleotide sequence of the endogenous gene. The transgene may also be selectively introduced into a particular cell type, thus inactivating the endogenous gene in only that cell type, by following, for example, the teaching of Gu et al. (Gu et al., Science 265:103-106 (1994)). The regulatory sequences required for such a cell-type specific inactivation will depend upon the particular cell type of interest, and will be apparent to those of skill in the art.

[1011] Once transgenic animals have been generated, the expression of the recombinant gene may be assayed utilizing standard techniques. Initial screening may be accomplished by Southern blot analysis or PCR techniques to analyze animal tissues to verify that integration of the transgene has taken place. The level of mRNA expression of the transgene in the tissues of the transgenic animals may also be assessed using techniques which include, but are not limited to, Northern blot analysis of tissue samples obtained from the animal, *in situ* hybridization analysis, and reverse transcriptase-PCR (rt-PCR). Samples of transgenic gene-expressing tissue may also be evaluated immunocytochemically or immunohistochemically using antibodies specific for the transgene product.

[1012] Once the founder animals are produced, they may be bred, inbred, outbred, or crossbred to produce colonies of the particular animal. Examples of such breeding strategies include, but are not limited to: outbreeding of founder animals with more than one integration site in order to establish separate lines; inbreeding of separate lines in order to produce compound transgenics that express the transgene at higher levels because of the effects of additive expression of each transgene; crossing of heterozygous

transgenic animals to produce animals homozygous for a given integration site in order to both augment expression and eliminate the need for screening of animals by DNA analysis; crossing of separate homozygous lines to produce compound heterozygous or homozygous lines; and breeding to place the transgene on a distinct background that is appropriate for an experimental model of interest.

[1013] Transgenic animals of the invention have uses which include, but are not limited to, animal model systems useful in elaborating the biological function of polypeptides of the present invention, studying conditions and/or disorders associated with aberrant expression, and in screening for compounds effective in ameliorating such conditions and/or disorders.

Example 20: Knock-Out Animals

[1014] Endogenous gene expression can also be reduced by inactivating or "knocking out" the gene and/or its promoter using targeted homologous recombination. (e.g., see Smithies et al., *Nature* 317:230-234 (1985); Thomas & Capecchi, *Cell* 51:503-512 (1987); Thompson et al., *Cell* 5:313-321 (1989); each of which is incorporated by reference herein in its entirety). For example, a mutant, non-functional polynucleotide of the invention (or a completely unrelated DNA sequence) flanked by DNA homologous to the endogenous polynucleotide sequence (either the coding regions or regulatory regions of the gene) can be used, with or without a selectable marker and/or a negative selectable marker, to transfect cells that express polypeptides of the invention *in vivo*. In another embodiment, techniques known in the art are used to generate knockouts in cells that contain, but do not express the gene of interest. Insertion of the DNA construct, via targeted homologous recombination, results in inactivation of the targeted gene. Such approaches are particularly suited in research and agricultural fields where modifications to embryonic stem cells can be used to generate animal offspring with an inactive targeted gene (e.g., see Thomas & Capecchi 1987 and Thompson 1989, *supra*). However this approach can be routinely adapted for use in humans provided the recombinant DNA constructs are directly administered or targeted to the required site *in vivo* using appropriate viral vectors that will be apparent to those of skill in the art.

[1015] In further embodiments of the invention, cells that are genetically engineered to express the polypeptides of the invention, or alternatively, that are genetically engineered not to express the polypeptides of the invention (e.g., knockouts) are administered to a patient *in vivo*. Such cells may be obtained from the patient (i.e., animal, including human) or an MHC compatible donor and can include, but are not limited to fibroblasts, bone marrow cells, blood cells (e.g., lymphocytes), adipocytes, muscle cells, endothelial cells etc. The cells are genetically engineered *in vitro* using recombinant DNA techniques to introduce the coding sequence of polypeptides of the invention into the cells, or alternatively, to disrupt the coding sequence and/or endogenous regulatory sequence associated with the polypeptides of the invention, e.g., by transduction (using viral vectors, and preferably vectors that integrate the transgene into the cell genome) or transfection procedures, including, but not limited to, the use of plasmids, cosmids, YACs, naked DNA, electroporation, liposomes, etc. The coding sequence of the polypeptides of the invention can be placed under the control of a strong constitutive or inducible promoter or promoter/enhancer to achieve expression, and preferably secretion, of the polypeptides of the invention. The engineered cells which express and preferably secrete the polypeptides of the invention can be introduced into the patient systemically, e.g., in the circulation, or intraperitoneally.

[1016] Alternatively, the cells can be incorporated into a matrix and implanted in the body, e.g., genetically engineered fibroblasts can be implanted as part of a skin graft; genetically engineered endothelial cells can be implanted as part of a lymphatic or vascular graft. (See, for example, Anderson et al. U.S. Patent No. 5,399,349; and Mulligan & Wilson, U.S. Patent No. 5,460,959 each of which is incorporated by reference herein in its entirety).

[1017] When the cells to be administered are non-autologous or non-MHC compatible cells, they can be administered using well known techniques which prevent the development of a host immune response against the introduced cells. For example, the cells may be introduced in an encapsulated form which, while allowing for an exchange of components with the immediate extracellular environment, does not allow the introduced cells to be recognized by the host immune system.

[1018] Transgenic and "knock-out" animals of the invention have uses which include, but are not limited to, animal model systems useful in elaborating the biological function

of polypeptides of the present invention, studying conditions and/or disorders associated with aberrant expression, and in screening for compounds effective in ameliorating such conditions and/or disorders.

***Example 21: Assays Detecting Stimulation or Inhibition of B cell
Proliferation and Differentiation***

[1019] Generation of functional humoral immune responses requires both soluble and cognate signaling between B-lineage cells and their microenvironment. Signals may impart a positive stimulus that allows a B-lineage cell to continue its programmed development, or a negative stimulus that instructs the cell to arrest its current developmental pathway. To date, numerous stimulatory and inhibitory signals have been found to influence B cell responsiveness including IL-2, IL-4, IL-5, IL-6, IL-7, IL10, IL-13, IL-14 and IL-15. Interestingly, these signals are by themselves weak effectors but can, in combination with various co-stimulatory proteins, induce activation, proliferation, differentiation, homing, tolerance and death among B cell populations.

[1020] One of the best studied classes of B-cell co-stimulatory proteins is the TNF-superfamily. Within this family CD40, CD27, and CD30 along with their respective ligands CD154, CD70, and CD153 have been found to regulate a variety of immune responses. Assays which allow for the detection and/or observation of the proliferation and differentiation of these B-cell populations and their precursors are valuable tools in determining the effects various proteins may have on these B-cell populations in terms of proliferation and differentiation. Listed below are two assays designed to allow for the detection of the differentiation, proliferation, or inhibition of B-cell populations and their precursors.

[1021] *In Vitro* Assay- Agonists or antagonists of the invention can be assessed for its ability to induce activation, proliferation, differentiation or inhibition and/or death in B-cell populations and their precursors. The activity of the agonists or antagonists of the invention on purified human tonsillar B cells, measured qualitatively over the dose range from 0.1 to 10,000 ng/mL, is assessed in a standard B-lymphocyte co-stimulation assay in which purified tonsillar B cells are cultured in the presence of either formalin-fixed *Staphylococcus aureus* Cowan I (SAC) or immobilized anti-human IgM antibody as the

priming agent. Second signals such as IL-2 and IL-15 synergize with SAC and IgM crosslinking to elicit B cell proliferation as measured by tritiated-thymidine incorporation. Novel synergizing agents can be readily identified using this assay. The assay involves isolating human tonsillar B cells by magnetic bead (MACS) depletion of CD3-positive cells. The resulting cell population is greater than 95% B cells as assessed by expression of CD45R(B220).

[1022] Various dilutions of each sample are placed into individual wells of a 96-well plate to which are added 10^5 B-cells suspended in culture medium (RPMI 1640 containing 10% FBS, 5×10^{-5} M 2ME, 100U/ml penicillin, 10ug/ml streptomycin, and 10^{-5} dilution of SAC) in a total volume of 150ul. Proliferation or inhibition is quantitated by a 20h pulse (1uCi/well) with 3 H-thymidine (6.7 Ci/mM) beginning 72h post factor addition. The positive and negative controls are IL2 and medium respectively.

[1023] *In vivo* Assay- BALB/c mice are injected (i.p.) twice per day with buffer only, or 2 mg/Kg of agonists or antagonists of the invention, or truncated forms thereof. Mice receive this treatment for 4 consecutive days, at which time they are sacrificed and various tissues and serum collected for analyses. Comparison of H&E sections from normal spleens and spleens treated with agonists or antagonists of the invention identify the results of the activity of the agonists or antagonists on spleen cells, such as the diffusion of peri-arterial lymphatic sheaths, and/or significant increases in the nucleated cellularity of the red pulp regions, which may indicate the activation of the differentiation and proliferation of B-cell populations. Immunohistochemical studies using a B cell marker, anti-CD45R(B220), are used to determine whether any physiological changes to splenic cells, such as splenic disorganization, are due to increased B-cell representation within loosely defined B-cell zones that infiltrate established T-cell regions.

[1024] Flow cytometric analyses of the spleens from mice treated with agonist or antagonist is used to indicate whether the agonists or antagonists specifically increases the proportion of ThB+, CD45R(B220)dull B cells over that which is observed in control mice.

[1025] Likewise, a predicted consequence of increased mature B-cell representation *in vivo* is a relative increase in serum Ig titers. Accordingly, serum IgM and IgA levels are compared between buffer and agonists or antagonists-treated mice.

[1026] The studies described in this example tested activity of agonists or antagonists of the invention. However, one skilled in the art could easily modify the exemplified studies to test the activity of polynucleotides or polypeptides of the invention (e.g., gene therapy).

Example 22: T Cell Proliferation Assay

[1027] A CD3-induced proliferation assay is performed on PBMCs and is measured by the uptake of ^3H -thymidine. The assay is performed as follows. Ninety-six well plates are coated with 100 μl /well of mAb to CD3 (HIT3a, Pharmingen) or isotype-matched control mAb (B33.1) overnight at 4 degrees C (1 $\mu\text{g}/\text{ml}$ in .05M bicarbonate buffer, pH 9.5), then washed three times with PBS. PBMC are isolated by F/H gradient centrifugation from human peripheral blood and added to quadruplicate wells (5×10^4 /well) of mAb coated plates in RPMI containing 10% FCS and P/S in the presence of varying concentrations of agonists or antagonists of the invention (total volume 200 μl). Relevant protein buffer and medium alone are controls. After 48 hr. culture at 37 degrees C, plates are spun for 2 min. at 1000 rpm and 100 μl of supernatant is removed and stored -20 degrees C for measurement of IL-2 (or other cytokines) if effect on proliferation is observed. Wells are supplemented with 100 μl of medium containing 0.5 uCi of ^3H -thymidine and cultured at 37 degrees C for 18-24 hr. Wells are harvested and incorporation of ^3H -thymidine used as a measure of proliferation. Anti-CD3 alone is the positive control for proliferation. IL-2 (100 U/ml) is also used as a control which enhances proliferation. Control antibody which does not induce proliferation of T cells is used as the negative control for the effects of agonists or antagonists of the invention.

[1028] The studies described in this example tested activity of agonists or antagonists of the invention. However, one skilled in the art could easily modify the exemplified studies to test the activity of polynucleotides or polypeptides of the invention (e.g., gene therapy).

Example 23: Effect of Agonists or Antagonists of the Invention on the Expression of MHC Class II, Costimulatory and Adhesion Molecules and Cell Differentiation of Monocytes and Monocyte-Derived Human Dendritic Cells

[1029] Dendritic cells are generated by the expansion of proliferating precursors found in the peripheral blood: adherent PBMC or elutriated monocytic fractions are cultured for 7-14 days with GM-CSF (50 ng/ml) and IL-4 (20 ng/ml). These dendritic cells have the characteristic phenotype of immature cells (expression of CD1, CD80, CD86, CD40 and MHC class II antigens). Treatment with activating factors, such as TNF- α , causes a rapid change in surface phenotype (increased expression of MHC class I and II, costimulatory and adhesion molecules, downregulation of FC γ RII, upregulation of CD83). These changes correlate with increased antigen-presenting capacity and with functional maturation of the dendritic cells.

[1030] FACS analysis of surface antigens is performed as follows. Cells are treated 1-4 days with increasing concentrations of agonist or antagonist of the invention or LPS (positive control), washed with PBS containing 1% BSA and 0.02 mM sodium azide, and then incubated with 1:20 dilution of appropriate FITC- or PE-labeled monoclonal antibodies for 30 minutes at 4 degrees C. After an additional wash, the labeled cells are analyzed by flow cytometry on a FACScan (Becton Dickinson).

[1031] Effect on the production of cytokines. Cytokines generated by dendritic cells, in particular IL-12, are important in the initiation of T-cell dependent immune responses. IL-12 strongly influences the development of Th1 helper T-cell immune response, and induces cytotoxic T and NK cell function. An ELISA is used to measure the IL-12 release as follows. Dendritic cells (10^6 /ml) are treated with increasing concentrations of agonists or antagonists of the invention for 24 hours. LPS (100 ng/ml) is added to the cell culture as positive control. Supernatants from the cell cultures are then collected and analyzed for IL-12 content using commercial ELISA kit (e.g., R & D Systems (Minneapolis, MN)). The standard protocols provided with the kits are used.

[1032] Effect on the expression of MHC Class II, costimulatory and adhesion molecules. Three major families of cell surface antigens can be identified on monocytes: adhesion molecules, molecules involved in antigen presentation, and Fc receptor. Modulation of the expression of MHC class II antigens and other costimulatory molecules, such as B7 and ICAM-1, may result in changes in the antigen presenting capacity of monocytes and ability to induce T cell activation. Increased expression of Fc receptors may correlate with improved monocyte cytotoxic activity, cytokine release and phagocytosis.

[1033] FACS analysis is used to examine the surface antigens as follows. Monocytes are treated 1-5 days with increasing concentrations of agonists or antagonists of the invention or LPS (positive control), washed with PBS containing 1% BSA and 0.02 mM sodium azide, and then incubated with 1:20 dilution of appropriate FITC- or PE-labeled monoclonal antibodies for 30 minutes at 4 degrees C. After an additional wash, the labeled cells are analyzed by flow cytometry on a FACScan (Becton Dickinson).

[1034] Monocyte activation and/or increased survival. Assays for molecules that activate (or alternatively, inactivate) monocytes and/or increase monocyte survival (or alternatively, decrease monocyte survival) are known in the art and may routinely be applied to determine whether a molecule of the invention functions as an inhibitor or activator of monocytes. Agonists or antagonists of the invention can be screened using the three assays described below. For each of these assays, Peripheral blood mononuclear cells (PBMC) are purified from single donor leukopacks (American Red Cross, Baltimore, MD) by centrifugation through a Histopaque gradient (Sigma). Monocytes are isolated from PBMC by counterflow centrifugal elutriation.

[1035] Monocyte Survival Assay. Human peripheral blood monocytes progressively lose viability when cultured in absence of serum or other stimuli. Their death results from internally regulated processes (apoptosis). Addition to the culture of activating factors, such as TNF-alpha dramatically improves cell survival and prevents DNA fragmentation. Propidium iodide (PI) staining is used to measure apoptosis as follows. Monocytes are cultured for 48 hours in polypropylene tubes in serum-free medium (positive control), in the presence of 100 ng/ml TNF-alpha (negative control), and in the presence of varying

concentrations of the compound to be tested. Cells are suspended at a concentration of 2×10^6 /ml in PBS containing PI at a final concentration of 5 μ g/ml, and then incubated at room temperature for 5 minutes before FACScan analysis. PI uptake has been demonstrated to correlate with DNA fragmentation in this experimental paradigm.

[1036] Effect on cytokine release. An important function of monocytes/macrophages is their regulatory activity on other cellular populations of the immune system through the release of cytokines after stimulation. An ELISA to measure cytokine release is performed as follows. Human monocytes are incubated at a density of 5×10^5 cells/ml with increasing concentrations of agonists or antagonists of the invention and under the same conditions, but in the absence of agonists or antagonists. For IL-12 production, the cells are primed overnight with IFN (100 U/ml) in the presence of agonist or antagonist of the invention. LPS (10 ng/ml) is then added. Conditioned media are collected after 24h and kept frozen until use. Measurement of TNF-alpha, IL-10, MCP-1 and IL-8 is then performed using a commercially available ELISA kit (e.g., R & D Systems (Minneapolis, MN)) and applying the standard protocols provided with the kit.

[1037] Oxidative burst. Purified monocytes are plated in 96-w plate at 2×10^5 cell/well. Increasing concentrations of agonists or antagonists of the invention are added to the wells in a total volume of 0.2 ml culture medium (RPMI 1640 + 10% FCS, glutamine and antibiotics). After 3 days incubation, the plates are centrifuged and the medium is removed from the wells. To the macrophage monolayers, 0.2 ml per well of phenol red solution (140 mM NaCl, 10 mM potassium phosphate buffer pH 7.0, 5.5 mM dextrose, 0.56 mM phenol red and 19 U/ml of HRPO) is added, together with the stimulant (200 nM PMA). The plates are incubated at 37°C for 2 hours and the reaction is stopped by adding 20 μ l 1N NaOH per well. The absorbance is read at 610 nm. To calculate the amount of H_2O_2 produced by the macrophages, a standard curve of a H_2O_2 solution of known molarity is performed for each experiment.

[1038] The studies described in this example tested activity of agonists or antagonists of the invention. However, one skilled in the art could easily modify the exemplified studies to test the activity of polynucleotides or polypeptides of the invention (e.g., gene therapy).

Example 24: Biological Effects of Agonists or Antagonists of the Invention**Astrocyte and Neuronal Assays:**

[1039] Agonists or antagonists of the invention, expressed in *Escherichia coli* and purified as described above, can be tested for activity in promoting the survival, neurite outgrowth, or phenotypic differentiation of cortical neuronal cells and for inducing the proliferation of glial fibrillary acidic protein immunopositive cells, astrocytes. The selection of cortical cells for the bioassay is based on the prevalent expression of FGF-1 and FGF-2 in cortical structures and on the previously reported enhancement of cortical neuronal survival resulting from FGF-2 treatment. A thymidine incorporation assay, for example, can be used to elucidate an agonist or antagonist of the invention's activity on these cells.

[1040] Moreover, previous reports describing the biological effects of FGF-2 (basic FGF) on cortical or hippocampal neurons *in vitro* have demonstrated increases in both neuron survival and neurite outgrowth (Walicke et al., "Fibroblast growth factor promotes survival of dissociated hippocampal neurons and enhances neurite extension." *Proc. Natl. Acad. Sci. USA* 83:3012-3016. (1986), assay herein incorporated by reference in its entirety). However, reports from experiments done on PC-12 cells suggest that these two responses are not necessarily synonymous and may depend on not only which FGF is being tested but also on which receptor(s) are expressed on the target cells. Using the primary cortical neuronal culture paradigm, the ability of an agonist or antagonist of the invention to induce neurite outgrowth can be compared to the response achieved with FGF-2 using, for example, a thymidine incorporation assay.

Fibroblast and endothelial cell assays:

[1041] Human lung fibroblasts are obtained from Clonetics (San Diego, CA) and maintained in growth media from Clonetics. Dermal microvascular endothelial cells are obtained from Cell Applications (San Diego, CA). For proliferation assays, the human lung fibroblasts and dermal microvascular endothelial cells can be cultured at 5,000 cells/well in a 96-well plate for one day in growth medium. The cells are then incubated for one day in 0.1% BSA basal medium. After replacing the medium with fresh 0.1% BSA medium, the cells are

incubated with the test proteins for 3 days. Alamar Blue (Alamar Biosciences, Sacramento, CA) is added to each well to a final concentration of 10%. The cells are incubated for 4 hr. Cell viability is measured by reading in a CytoFluor fluorescence reader. For the PGE₂ assays, the human lung fibroblasts are cultured at 5,000 cells/well in a 96-well plate for one day. After a medium change to 0.1% BSA basal medium, the cells are incubated with FGF-2 or agonists or antagonists of the invention with or without IL-1 α for 24 hours. The supernatants are collected and assayed for PGE₂ by EIA kit (Cayman, Ann Arbor, MI). For the IL-6 assays, the human lung fibroblasts are cultured at 5,000 cells/well in a 96-well plate for one day. After a medium change to 0.1% BSA basal medium, the cells are incubated with FGF-2 or with or without agonists or antagonists of the invention IL-1 α for 24 hours. The supernatants are collected and assayed for IL-6 by ELISA kit (Endogen, Cambridge, MA).

[1042] Human lung fibroblasts are cultured with FGF-2 or agonists or antagonists of the invention for 3 days in basal medium before the addition of Alamar Blue to assess effects on growth of the fibroblasts. FGF-2 should show a stimulation at 10 - 2500 ng/ml which can be used to compare stimulation with agonists or antagonists of the invention.

Parkinson Models.

[1043] The loss of motor function in Parkinson's disease is attributed to a deficiency of striatal dopamine resulting from the degeneration of the nigrostriatal dopaminergic projection neurons. An animal model for Parkinson's that has been extensively characterized involves the systemic administration of 1-methyl-4 phenyl 1,2,3,6-tetrahydropyridine (MPTP). In the CNS, MPTP is taken-up by astrocytes and catabolized by monoamine oxidase B to 1-methyl-4-phenyl pyridine (MPP⁺) and released. Subsequently, MPP⁺ is actively accumulated in dopaminergic neurons by the high-affinity reuptake transporter for dopamine. MPP⁺ is then concentrated in mitochondria by the electrochemical gradient and selectively inhibits nicotinamide adenine disphosphate: ubiquinone oxidoreductionase (complex I), thereby interfering with electron transport and eventually generating oxygen radicals.

[1044] It has been demonstrated in tissue culture paradigms that FGF-2 (basic FGF) has trophic activity towards nigral dopaminergic neurons (Ferrari et al., Dev. Biol. 1989). Recently, Dr. Unsicker's group has demonstrated that administering FGF-2 in gel foam

implants in the striatum results in the near complete protection of nigral dopaminergic neurons from the toxicity associated with MPTP exposure (Otto and Unsicker, J. Neuroscience, 1990).

[1045] Based on the data with FGF-2, agonists or antagonists of the invention can be evaluated to determine whether it has an action similar to that of FGF-2 in enhancing dopaminergic neuronal survival *in vitro* and it can also be tested *in vivo* for protection of dopaminergic neurons in the striatum from the damage associated with MPTP treatment. The potential effect of an agonist or antagonist of the invention is first examined *in vitro* in a dopaminergic neuronal cell culture paradigm. The cultures are prepared by dissecting the midbrain floor plate from gestation day 14 Wistar rat embryos. The tissue is dissociated with trypsin and seeded at a density of 200,000 cells/cm² on polyorthinine-laminin coated glass coverslips. The cells are maintained in Dulbecco's Modified Eagle's medium and F12 medium containing hormonal supplements (N1). The cultures are fixed with paraformaldehyde after 8 days *in vitro* and are processed for tyrosine hydroxylase, a specific marker for dopaminergic neurons, immunohistochemical staining. Dissociated cell cultures are prepared from embryonic rats. The culture medium is changed every third day and the factors are also added at that time.

[1046] Since the dopaminergic neurons are isolated from animals at gestation day 14, a developmental time which is past the stage when the dopaminergic precursor cells are proliferating, an increase in the number of tyrosine hydroxylase immunopositive neurons would represent an increase in the number of dopaminergic neurons surviving *in vitro*. Therefore, if an agonist or antagonist of the invention acts to prolong the survival of dopaminergic neurons, it would suggest that the agonist or antagonist may be involved in Parkinson's Disease.

[1047] The studies described in this example tested activity of agonists or antagonists of the invention. However, one skilled in the art could easily modify the exemplified studies to test the activity of polynucleotides or polypeptides of the invention (e.g., gene therapy).

Example 25: The Effect of Agonists or Antagonists of the Invention on the Growth of Vascular Endothelial Cells

[1048] On day 1, human umbilical vein endothelial cells (HUVEC) are seeded at $2-5 \times 10^4$ cells/35 mm dish density in M199 medium containing 4% fetal bovine serum (FBS), 16 units/ml heparin, and 50 units/ml endothelial cell growth supplements (ECGS, Biotechnology, Inc.). On day 2, the medium is replaced with M199 containing 10% FBS, 8 units/ml heparin. An agonist or antagonist of the invention, and positive controls, such as VEGF and basic FGF (bFGF) are added, at varying concentrations. On days 4 and 6, the medium is replaced. On day 8, cell number is determined with a Coulter Counter.

[1049] An increase in the number of HUVEC cells indicates that the compound of the invention may proliferate vascular endothelial cells, while a decrease in the number of HUVEC cells indicates that the compound of the invention inhibits vascular endothelial cells.

[1050] The studies described in this example tested activity of a polypeptide of the invention. However, one skilled in the art could easily modify the exemplified studies to test the activity of polynucleotides (e.g., gene therapy), agonists, and/or antagonists of the invention.

Example 26: Rat Corneal Wound Healing Model

[1051] This animal model shows the effect of an agonist or antagonist of the invention on neovascularization. The experimental protocol includes:

Making a 1-1.5 mm long incision from the center of cornea into the stromal layer.

Inserting a spatula below the lip of the incision facing the outer corner of the eye.

Making a pocket (its base is 1-1.5 mm from the edge of the eye).

Positioning a pellet, containing 50ng- 5ug of an agonist or antagonist of the invention, within the pocket.

[1052] Treatment with an agonist or antagonist of the invention can also be applied topically to the corneal wounds in a dosage range of 20mg - 500mg (daily treatment for five days).

[1053] The studies described in this example tested activity of agonists or antagonists of the invention. However, one skilled in the art could easily modify the exemplified studies to test the activity of polynucleotides or polypeptides of the invention (e.g., gene therapy).

***Example 27: Diabetic Mouse and Glucocorticoid-Impaired
Wound Healing Models***

[1054] *Diabetic db+/db+ Mouse Model.*

[1055] To demonstrate that an agonist or antagonist of the invention accelerates the healing process, the genetically diabetic mouse model of wound healing is used. The full thickness wound healing model in the db+/db+ mouse is a well characterized, clinically relevant and reproducible model of impaired wound healing. Healing of the diabetic wound is dependent on formation of granulation tissue and re-epithelialization rather than contraction (Gartner, M.H. *et al.*, *J. Surg. Res.* 52:389 (1992); Greenhalgh, D.G. *et al.*, *Am. J. Pathol.* 136:1235 (1990)).

[1056] The diabetic animals have many of the characteristic features observed in Type II diabetes mellitus. Homozygous (db+/db+) mice are obese in comparison to their normal heterozygous (db+/+m) littermates. Mutant diabetic (db+/db+) mice have a single autosomal recessive mutation on chromosome 4 (db+) (Coleman *et al.* *Proc. Natl. Acad. Sci. USA* 77:283-293 (1982)). Animals show polyphagia, polydipsia and polyuria. Mutant diabetic mice (db+/db+) have elevated blood glucose, increased or normal insulin levels, and suppressed cell-mediated immunity (Mandel *et al.*, *J. Immunol.* 120:1375 (1978); Debray-Sachs, M. *et al.*, *Clin. Exp. Immunol.* 51(1):1-7 (1983); Leiter *et al.*, *Am. J. of Pathol.* 114:46-55 (1985)). Peripheral neuropathy, myocardial complications, and microvascular lesions, basement membrane thickening and glomerular filtration abnormalities have been described in these animals (Norido, F. *et al.*, *Exp. Neurol.* 83(2):221-232 (1984); Robertson *et al.*, *Diabetes* 29(1):60-67 (1980); Giacomelli *et al.*, *Lab Invest.* 40(4):460-473 (1979); Coleman, D.L., *Diabetes* 31 (Suppl):1-6 (1982)). These homozygous diabetic mice develop hyperglycemia that is resistant to insulin analogous to human type II diabetes (Mandel *et al.*, *J. Immunol.* 120:1375-1377 (1978)).

[1057] The characteristics observed in these animals suggests that healing in this model may be similar to the healing observed in human diabetes (Greenhalgh, *et al.*, *Am. J. of Pathol.* 136:1235-1246 (1990)).

[1058] Genetically diabetic female C57BL/KsJ (db+/db+) mice and their non-diabetic (db+/+m) heterozygous littermates are used in this study (Jackson Laboratories). The animals are purchased at 6 weeks of age and are 8 weeks old at the beginning of the study. Animals

are individually housed and received food and water ad libitum. All manipulations are performed using aseptic techniques. The experiments are conducted according to the rules and guidelines of Human Genome Sciences, Inc. Institutional Animal Care and Use Committee and the Guidelines for the Care and Use of Laboratory Animals.

[1059] Wounding protocol is performed according to previously reported methods (Tsuboi, R. and Rifkin, D.B., *J. Exp. Med.* 172:245-251 (1990)). Briefly, on the day of wounding, animals are anesthetized with an intraperitoneal injection of Avertin (0.01 mg/mL), 2,2,2-tribromoethanol and 2-methyl-2-butanol dissolved in deionized water. The dorsal region of the animal is shaved and the skin washed with 70% ethanol solution and iodine. The surgical area is dried with sterile gauze prior to wounding. An 8 mm full-thickness wound is then created using a Keyes tissue punch. Immediately following wounding, the surrounding skin is gently stretched to eliminate wound expansion. The wounds are left open for the duration of the experiment. Application of the treatment is given topically for 5 consecutive days commencing on the day of wounding. Prior to treatment, wounds are gently cleansed with sterile saline and gauze sponges.

[1060] Wounds are visually examined and photographed at a fixed distance at the day of surgery and at two day intervals thereafter. Wound closure is determined by daily measurement on days 1-5 and on day 8. Wounds are measured horizontally and vertically using a calibrated Jameson caliper. Wounds are considered healed if granulation tissue is no longer visible and the wound is covered by a continuous epithelium.

[1061] An agonist or antagonist of the invention is administered using at a range different doses, from 4mg to 500mg per wound per day for 8 days in vehicle. Vehicle control groups received 50mL of vehicle solution.

[1062] Animals are euthanized on day 8 with an intraperitoneal injection of sodium pentobarbital (300mg/kg). The wounds and surrounding skin are then harvested for histology and immunohistochemistry. Tissue specimens are placed in 10% neutral buffered formalin in tissue cassettes between biopsy sponges for further processing.

[1063] Three groups of 10 animals each (5 diabetic and 5 non-diabetic controls) are evaluated: 1) Vehicle placebo control, 2) untreated group, and 3) treated group.

[1064] Wound closure is analyzed by measuring the area in the vertical and horizontal axis and obtaining the total square area of the wound. Contraction is then estimated by establishing the differences between the initial wound area (day 0) and that of post treatment (day 8). The

wound area on day 1 is 64mm^2 , the corresponding size of the dermal punch. Calculations are made using the following formula:

$$[\text{Open area on day 8}] - [\text{Open area on day 1}] / [\text{Open area on day 1}]$$

[1065] Specimens are fixed in 10% buffered formalin and paraffin embedded blocks are sectioned perpendicular to the wound surface (5mm) and cut using a Reichert-Jung microtome. Routine hematoxylin-eosin (H&E) staining is performed on cross-sections of bisected wounds. Histologic examination of the wounds are used to assess whether the healing process and the morphologic appearance of the repaired skin is altered by treatment with an agonist or antagonist of the invention. This assessment included verification of the presence of cell accumulation, inflammatory cells, capillaries, fibroblasts, re-epithelialization and epidermal maturity (Greenhalgh, D.G. *et al.*, *Am. J. Pathol.* 136:1235 (1990)). A calibrated lens micrometer is used by a blinded observer.

[1066] Tissue sections are also stained immunohistochemically with a polyclonal rabbit anti-human keratin antibody using ABC Elite detection system. Human skin is used as a positive tissue control while non-immune IgG is used as a negative control. Keratinocyte growth is determined by evaluating the extent of reepithelialization of the wound using a calibrated lens micrometer.

[1067] Proliferating cell nuclear antigen/cyclin (PCNA) in skin specimens is demonstrated by using anti-PCNA antibody (1:50) with an ABC Elite detection system. Human colon cancer served as a positive tissue control and human brain tissue is used as a negative tissue control. Each specimen included a section with omission of the primary antibody and substitution with non-immune mouse IgG. Ranking of these sections is based on the extent of proliferation on a scale of 0-8, the lower side of the scale reflecting slight proliferation to the higher side reflecting intense proliferation.

[1068] Experimental data are analyzed using an unpaired t test. A p value of < 0.05 is considered significant.

[1069] *Steroid Impaired Rat Model*

[1070] The inhibition of wound healing by steroids has been well documented in various *in vitro* and *in vivo* systems (Wahl, Glucocorticoids and Wound healing. In: Anti-Inflammatory

Steroid Action: Basic and Clinical Aspects. 280-302 (1989); Wahlet *et al.*, *J. Immunol.* 115: 476-481 (1975); Werb *et al.*, *J. Exp. Med.* 147:1684-1694 (1978)). Glucocorticoids retard wound healing by inhibiting angiogenesis, decreasing vascular permeability (Ebert *et al.*, *An. Intern. Med.* 37:701-705 (1952)), fibroblast proliferation, and collagen synthesis (Beck *et al.*, *Growth Factors.* 5: 295-304 (1991); Haynes *et al.*, *J. Clin. Invest.* 61: 703-797 (1978)) and producing a transient reduction of circulating monocytes (Haynes *et al.*, *J. Clin. Invest.* 61: 703-797 (1978); Wahl, "Glucocorticoids and wound healing", *In: Antiinflammatory Steroid Action: Basic and Clinical Aspects*, Academic Press, New York, pp. 280-302 (1989)). The systemic administration of steroids to impaired wound healing is a well establish phenomenon in rats (Beck *et al.*, *Growth Factors.* 5: 295-304 (1991); Haynes *et al.*, *J. Clin. Invest.* 61: 703-797 (1978); Wahl, "Glucocorticoids and wound healing", *In: Antiinflammatory Steroid Action: Basic and Clinical Aspects*, Academic Press, New York, pp. 280-302 (1989); Pierce *et al.*, *Proc. Natl. Acad. Sci. USA* 86: 2229-2233 (1989)).

[1071] To demonstrate that an agonist or antagonist of the invention can accelerate the healing process, the effects of multiple topical applications of the agonist or antagonist on full thickness excisional skin wounds in rats in which healing has been impaired by the systemic administration of methylprednisolone is assessed.

[1072] Young adult male Sprague Dawley rats weighing 250-300 g (Charles River Laboratories) are used in this example. The animals are purchased at 8 weeks of age and are 9 weeks old at the beginning of the study. The healing response of rats is impaired by the systemic administration of methylprednisolone (17mg/kg/rat intramuscularly) at the time of wounding. Animals are individually housed and received food and water *ad libitum*. All manipulations are performed using aseptic techniques. This study is conducted according to the rules and guidelines of Human Genome Sciences, Inc. Institutional Animal Care and Use Committee and the Guidelines for the Care and Use of Laboratory Animals.

[1073] The wounding protocol is followed according to section A, above. On the day of wounding, animals are anesthetized with an intramuscular injection of ketamine (50 mg/kg) and xylazine (5 mg/kg). The dorsal region of the animal is shaved and the skin washed with 70% ethanol and iodine solutions. The surgical area is dried with sterile gauze prior to wounding. An 8 mm full-thickness wound is created using a Keyes tissue punch. The wounds are left open for the duration of the experiment. Applications of the testing materials are given topically once a day for 7 consecutive days commencing on the day of wounding and

subsequent to methylprednisolone administration. Prior to treatment, wounds are gently cleansed with sterile saline and gauze sponges.

[1074] Wounds are visually examined and photographed at a fixed distance at the day of wounding and at the end of treatment. Wound closure is determined by daily measurement on days 1-5 and on day 8. Wounds are measured horizontally and vertically using a calibrated Jameson caliper. Wounds are considered healed if granulation tissue is no longer visible and the wound is covered by a continuous epithelium.

[1075] The agonist or antagonist of the invention is administered using at a range different doses, from 4mg to 500mg per wound per day for 8 days in vehicle. Vehicle control groups received 50mL of vehicle solution.

[1076] Animals are euthanized on day 8 with an intraperitoneal injection of sodium pentobarbital (300mg/kg). The wounds and surrounding skin are then harvested for histology. Tissue specimens are placed in 10% neutral buffered formalin in tissue cassettes between biopsy sponges for further processing.

[1077] Three groups of 10 animals each (5 with methylprednisolone and 5 without glucocorticoid) are evaluated: 1) Untreated group 2) Vehicle placebo control 3) treated groups.

[1078] Wound closure is analyzed by measuring the area in the vertical and horizontal axis and obtaining the total area of the wound. Closure is then estimated by establishing the differences between the initial wound area (day 0) and that of post treatment (day 8). The wound area on day 1 is 64mm², the corresponding size of the dermal punch. Calculations are made using the following formula:

$$[\text{Open area on day 8}] - [\text{Open area on day 1}] / [\text{Open area on day 1}]$$

[1079] Specimens are fixed in 10% buffered formalin and paraffin embedded blocks are sectioned perpendicular to the wound surface (5mm) and cut using an Olympus microtome. Routine hematoxylin-eosin (H&E) staining is performed on cross-sections of bisected wounds. Histologic examination of the wounds allows assessment of whether the healing process and the morphologic appearance of the repaired skin is improved by treatment with an agonist or antagonist of the invention. A calibrated lens micrometer is used by a blinded observer to determine the distance of the wound gap.

[1080] Experimental data are analyzed using an unpaired t test. A p value of < 0.05 is considered significant.

[1081] The studies described in this example tested activity of agonists or antagonists of the invention. However, one skilled in the art could easily modify the exemplified studies to test the activity of polynucleotides or polypeptides of the invention (e.g., gene therapy).

Example 28: Lymphadema Animal Model

[1082] The purpose of this experimental approach is to create an appropriate and consistent lymphedema model for testing the therapeutic effects of an agonist or antagonist of the invention in lymphangiogenesis and re-establishment of the lymphatic circulatory system in the rat hind limb. Effectiveness is measured by swelling volume of the affected limb, quantification of the amount of lymphatic vasculature, total blood plasma protein, and histopathology. Acute lymphedema is observed for 7-10 days. Perhaps more importantly, the chronic progress of the edema is followed for up to 3-4 weeks.

[1083] Prior to beginning surgery, blood sample is drawn for protein concentration analysis. Male rats weighing approximately ~350g are dosed with Pentobarbital. Subsequently, the right legs are shaved from knee to hip. The shaved area is swabbed with gauze soaked in 70% EtOH. Blood is drawn for serum total protein testing. Circumference and volumetric measurements are made prior to injecting dye into paws after marking 2 measurement levels (0.5 cm above heel, at mid-pt of dorsal paw). The intradermal dorsum of both right and left paws are injected with 0.05 ml of 1% Evan's Blue. Circumference and volumetric measurements are then made following injection of dye into paws.

[1084] Using the knee joint as a landmark, a mid-leg inguinal incision is made circumferentially allowing the femoral vessels to be located. Forceps and hemostats are used to dissect and separate the skin flaps. After locating the femoral vessels, the lymphatic vessel that runs along side and underneath the vessel(s) is located. The main lymphatic vessels in this area are then electrically coagulated or suture ligated.

[1085] Using a microscope, muscles in back of the leg (near the semitendinosus and adductors) are bluntly dissected. The popliteal lymph node is then located. The 2 proximal and 2 distal lymphatic vessels and distal blood supply of the popliteal node are then ligated by

suturing. The popliteal lymph node, and any accompanying adipose tissue, is then removed by cutting connective tissues.

[1086] Care is taken to control any mild bleeding resulting from this procedure. After lymphatics are occluded, the skin flaps are sealed by using liquid skin (Vetbond) (AJ Buck). The separated skin edges are sealed to the underlying muscle tissue while leaving a gap of ~0.5 cm around the leg. Skin also may be anchored by suturing to underlying muscle when necessary.

[1087] To avoid infection, animals are housed individually with mesh (no bedding). Recovering animals are checked daily through the optimal edematous peak, which typically occurred by day 5-7. The plateau edematous peak are then observed. To evaluate the intensity of the lymphedema, the circumference and volumes of 2 designated places on each paw before operation and daily for 7 days are measured. The effect of plasma proteins on lymphedema is determined and whether protein analysis is a useful testing perimeter is also investigated. The weights of both control and edematous limbs are evaluated at 2 places. Analysis is performed in a blind manner.

[1088] Circumference Measurements: Under brief gas anesthetic to prevent limb movement, a cloth tape is used to measure limb circumference. Measurements are done at the ankle bone and dorsal paw by 2 different people and those 2 readings are averaged. Readings are taken from both control and edematous limbs.

[1089] Volumetric Measurements: On the day of surgery, animals are anesthetized with Pentobarbital and are tested prior to surgery. For daily volumetrics animals are under brief halothane anesthetic (rapid immobilization and quick recovery), and both legs are shaved and equally marked using waterproof marker on legs. Legs are first dipped in water, then dipped into instrument to each marked level then measured by Buxco edema software(Chen/Victor). Data is recorded by one person, while the other is dipping the limb to marked area.

[1090] Blood-plasma protein measurements: Blood is drawn, spun, and serum separated prior to surgery and then at conclusion for total protein and Ca^{2+} comparison.

[1091] Limb Weight Comparison: After drawing blood, the animal is prepared for tissue collection. The limbs are amputated using a quillitine, then both experimental and control legs are cut at the ligature and weighed. A second weighing is done as the tibio-cacaneal joint is disarticulated and the foot is weighed.

[1092] Histological Preparations: The transverse muscle located behind the knee (popliteal) area is dissected and arranged in a metal mold, filled with freezeGel, dipped into cold methylbutane, placed into labeled sample bags at - 80EC until sectioning. Upon sectioning, the muscle is observed under fluorescent microscopy for lymphatics..

[1093] The studies described in this example tested activity of agonists or antagonists of the invention. However, one skilled in the art could easily modify the exemplified studies to test the activity of polynucleotides or polypeptides of the invention (e.g., gene therapy).

Example 29: Suppression of TNF alpha-induced adhesion molecule expression by an Agonist or Antagonist of the Invention

[1094] The recruitment of lymphocytes to areas of inflammation and angiogenesis involves specific receptor-ligand interactions between cell surface adhesion molecules (CAMs) on lymphocytes and the vascular endothelium. The adhesion process, in both normal and pathological settings, follows a multi-step cascade that involves intercellular adhesion molecule-1 (ICAM-1), vascular cell adhesion molecule-1 (VCAM-1), and endothelial leukocyte adhesion molecule-1 (E-selectin) expression on endothelial cells (EC). The expression of these molecules and others on the vascular endothelium determines the efficiency with which leukocytes may adhere to the local vasculature and extravasate into the local tissue during the development of an inflammatory response. The local concentration of cytokines and growth factor participate in the modulation of the expression of these CAMs.

[1095] Tumor necrosis factor alpha (TNF-a), a potent proinflammatory cytokine, is a stimulator of all three CAMs on endothelial cells and may be involved in a wide variety of inflammatory responses, often resulting in a pathological outcome.

[1096] The potential of an agonist or antagonist of the invention to mediate a suppression of TNF-a induced CAM expression can be examined. A modified ELISA assay which uses ECs as a solid phase absorbent is employed to measure the amount of CAM expression on TNF-a treated ECs when co-stimulated with a member of the FGF family of proteins.

[1097] To perform the experiment, human umbilical vein endothelial cell (HUVEC) cultures are obtained from pooled cord harvests and maintained in growth medium (EGM-2; Clonetics, San Diego, CA) supplemented with 10% FCS and 1% penicillin/streptomycin in a 37 degree C humidified incubator containing 5% CO₂. HUVECs are seeded in 96-well plates

at concentrations of 1×10^4 cells/well in EGM medium at 37 degree C for 18-24 hrs or until confluent. The monolayers are subsequently washed 3 times with a serum-free solution of RPMI-1640 supplemented with 100 U/ml penicillin and 100 mg/ml streptomycin, and treated with a given cytokine and/or growth factor(s) for 24 h at 37 degree C. Following incubation, the cells are then evaluated for CAM expression.

[1098] Human Umbilical Vein Endothelial cells (HUVECs) are grown in a standard 96 well plate to confluence. Growth medium is removed from the cells and replaced with 90 μ l of 199 Medium (10% FBS). Samples for testing and positive or negative controls are added to the plate in triplicate (in 10 μ l volumes). Plates are incubated at 37 degree C for either 5 h (selectin and integrin expression) or 24 h (integrin expression only). Plates are aspirated to remove medium and 100 μ l of 0.1% paraformaldehyde-PBS(with Ca^{++} and Mg^{++}) is added to each well. Plates are held at 4°C for 30 min.

[1099] Fixative is then removed from the wells and wells are washed 1X with PBS(+Ca,Mg)+0.5% BSA and drained. Do not allow the wells to dry. Add 10 μ l of diluted primary antibody to the test and control wells. Anti-ICAM-1-Biotin, Anti-VCAM-1-Biotin and Anti-E-selectin-Biotin are used at a concentration of 10 μ g/ml (1:10 dilution of 0.1 mg/ml stock antibody). Cells are incubated at 37°C for 30 min. in a humidified environment. Wells are washed X3 with PBS(+Ca,Mg)+0.5% BSA.

[1100] Then add 20 μ l of diluted ExtrAvidin-Alkaline Phosphatase (1:5,000 dilution) to each well and incubated at 37°C for 30 min. Wells are washed X3 with PBS(+Ca,Mg)+0.5% BSA. 1 tablet of p-Nitrophenol Phosphate pNPP is dissolved in 5 ml of glycine buffer (pH 10.4). 100 μ l of pNPP substrate in glycine buffer is added to each test well. Standard wells in triplicate are prepared from the working dilution of the ExtrAvidin-Alkaline Phosphatase in glycine buffer: 1:5,000 (10^0) > $10^{-0.5}$ > 10^{-1} > $10^{-1.5}$. 5 μ l of each dilution is added to triplicate wells and the resulting AP content in each well is 5.50 ng, 1.74 ng, 0.55 ng, 0.18 ng. 100 μ l of pNPP reagent must then be added to each of the standard wells. The plate must be incubated at 37°C for 4h. A volume of 50 μ l of 3M NaOH is added to all wells. The results are quantified on a plate reader at 405 nm. The background subtraction option is used on blank wells filled with glycine buffer only. The template is set up to indicate the concentration of AP-conjugate in each standard well [5.50 ng; 1.74 ng; 0.55 ng; 0.18 ng]. Results are indicated as amount of bound AP-conjugate in each sample.

[1101] The studies described in this example tested activity of agonists or antagonists of the invention. However, one skilled in the art could easily modify the exemplified studies to test the activity of polynucleotides or polypeptides of the invention (e.g., gene therapy).

Example 30: Production Of Polypeptide of the Invention For High-Throughput Screening Assays

[1102] The following protocol produces a supernatant containing polypeptide of the present invention to be tested. This supernatant can then be used in the Screening Assays described in Examples 32-41.

[1103] First, dilute Poly-D-Lysine (644 587 Boehringer-Mannheim) stock solution (1mg/ml in PBS) 1:20 in PBS (w/o calcium or magnesium 17-516F Biowhittaker) for a working solution of 50ug/ml. Add 200 ul of this solution to each well (24 well plates) and incubate at RT for 20 minutes. Be sure to distribute the solution over each well (note: a 12-channel pipetter may be used with tips on every other channel). Aspirate off the Poly-D-Lysine solution and rinse with 1ml PBS (Phosphate Buffered Saline). The PBS should remain in the well until just prior to plating the cells and plates may be poly-lysine coated in advance for up to two weeks.

[1104] Plate 293T cells (do not carry cells past P+20) at 2×10^5 cells/well in .5ml DMEM(Dulbecco's Modified Eagle Medium)(with 4.5 G/L glucose and L-glutamine (12-604F Biowhittaker))/10% heat inactivated FBS(14-503F Biowhittaker)/1x Penstrep(17-602E Biowhittaker). Let the cells grow overnight.

[1105] The next day, mix together in a sterile solution basin: 300 ul Lipofectamine (18324-012 Gibco/BRL) and 5ml Optimem I (31985070 Gibco/BRL)/96-well plate. With a small volume multi-channel pipetter, aliquot approximately 2ug of an expression vector containing a polynucleotide insert, produced by the methods described in Examples 8-10, into an appropriately labeled 96-well round bottom plate. With a multi-channel pipetter, add 50ul of the Lipofectamine/Optimem I mixture to each well. Pipette up and down gently to mix. Incubate at RT 15-45 minutes. After about 20 minutes, use a multi-channel pipetter to add 150ul Optimem I to each well. As a control, one plate of vector DNA lacking an insert should be transfected with each set of transfections.

[1106] Preferably, the transfection should be performed by tag-teaming the following tasks. By tag-teaming, hands on time is cut in half, and the cells do not spend too much time on PBS. First, person A aspirates off the media from four 24-well plates of cells, and then person B rinses each well with .5-1ml PBS. Person A then aspirates off PBS rinse, and person B, using a 12-channel pipetter with tips on every other channel, adds the 200ul of DNA/Lipofectamine/Optimem I complex to the odd wells first, then to the even wells, to each row on the 24-well plates. Incubate at 37 degree C for 6 hours.

[1107] While cells are incubating, prepare appropriate media, either 1%BSA in DMEM with 1x penstrep, or HGS CHO-5 media (116.6 mg/L of CaCl₂ (anhyd); 0.00130 mg/L CuSO₄·5H₂O; 0.050 mg/L of Fe(NO₃)₃·9H₂O; 0.417 mg/L of FeSO₄·7H₂O; 311.80 mg/L of KCl; 28.64 mg/L of MgCl₂; 48.84 mg/L of MgSO₄; 6995.50 mg/L of NaCl; 2400.0 mg/L of NaHCO₃; 62.50 mg/L of NaH₂PO₄·H₂O; 71.02 mg/L of Na₂HPO₄; 4320 mg/L of ZnSO₄·7H₂O; .002 mg/L of Arachidonic Acid ; 1.022 mg/L of Cholesterol; .070 mg/L of DL-alpha-Tocopherol-Acetate; 0.0520 mg/L of Linoleic Acid; 0.010 mg/L of Linolenic Acid; 0.010 mg/L of Myristic Acid; 0.010 mg/L of Oleic Acid; 0.010 mg/L of Palmitic Acid; 0.010 mg/L of Palmitic Acid; 100 mg/L of Pluronic F-68; 0.010 mg/L of Stearic Acid; 2.20 mg/L of Tween 80; 4551 mg/L of D-Glucose; 130.85 mg/ml of L- Alanine; 147.50 mg/ml of L-Arginine-HCL; 7.50 mg/ml of L-Asparagine-H₂O; 6.65 mg/ml of L-Aspartic Acid; 29.56 mg/ml of L-Cystine-2HCL-H₂O; 31.29 mg/ml of L-Cystine-2HCL; 7.35 mg/ml of L-Glutamic Acid; 365.0 mg/ml of L-Glutamine; 18.75 mg/ml of Glycine; 52.48 mg/ml of L-Histidine-HCL-H₂O; 106.97 mg/ml of L-Isoleucine; 111.45 mg/ml of L-Leucine; 163.75 mg/ml of L-Lysine HCL; 32.34 mg/ml of L-Methionine; 68.48 mg/ml of L-Phenylalanine; 40.0 mg/ml of L-Proline; 26.25 mg/ml of L-Serine; 101.05 mg/ml of L-Threonine; 19.22 mg/ml of L-Tryptophan; 91.79 mg/ml of L-Tyrosine-2Na-2H₂O; and 99.65 mg/ml of L-Valine; 0.0035 mg/L of Biotin; 3.24 mg/L of D-Ca Pantothenate; 11.78 mg/L of Choline Chloride; 4.65 mg/L of Folic Acid; 15.60 mg/L of i-Inositol; 3.02 mg/L of Niacinamide; 3.00 mg/L of Pyridoxal HCL; 0.031 mg/L of Pyridoxine HCL; 0.319 mg/L of Riboflavin; 3.17 mg/L of Thiamine HCL; 0.365 mg/L of Thymidine; 0.680 mg/L of Vitamin B₁₂; 25 mM of HEPES Buffer; 2.39 mg/L of Na Hypoxanthine; 0.105 mg/L of Lipoic Acid; 0.081 mg/L of Sodium Putrescine-2HCL; 55.0 mg/L of Sodium Pyruvate; 0.0067 mg/L of Sodium Selenite; 20uM of Ethanolamine;

0.122 mg/L of Ferric Citrate; 41.70 mg/L of Methyl-B-Cyclodextrin complexed with Linoleic Acid; 33.33 mg/L of Methyl-B-Cyclodextrin complexed with Oleic Acid; 10 mg/L of Methyl-B-Cyclodextrin complexed with Retinal Acetate. Adjust osmolarity to 327 mOsm) with 2mm glutamine and 1x penstrep. (BSA (81-068-3 Bayer) 100gm dissolved in 1L DMEM for a 10% BSA stock solution). Filter the media and collect 50 ul for endotoxin assay in 15ml polystyrene conical.

[1108] The transfection reaction is terminated, preferably by tag-teaming, at the end of the incubation period. Person A aspirates off the transfection media, while person B adds 1.5ml appropriate media to each well. Incubate at 37 degree C for 45 or 72 hours depending on the media used: 1%BSA for 45 hours or CHO-5 for 72 hours.

[1109] On day four, using a 300ul multichannel pipetter, aliquot 600ul in one 1ml deep well plate and the remaining supernatant into a 2ml deep well. The supernatants from each well can then be used in the assays described in Examples 32-39.

[1110] It is specifically understood that when activity is obtained in any of the assays described below using a supernatant, the activity originates from either the polypeptide of the present invention directly (e.g., as a secreted protein) or by polypeptide of the present invention inducing expression of other proteins, which are then secreted into the supernatant. Thus, the invention further provides a method of identifying the protein in the supernatant characterized by an activity in a particular assay.

Example 31: Construction of GAS Reporter Construct

[1111] One signal transduction pathway involved in the differentiation and proliferation of cells is called the Jaks-STATs pathway. Activated proteins in the Jaks-STATs pathway bind to gamma activation site "GAS" elements or interferon-sensitive responsive element ("ISRE"), located in the promoter of many genes. The binding of a protein to these elements alter the expression of the associated gene.

[1112] GAS and ISRE elements are recognized by a class of transcription factors called Signal Transducers and Activators of Transcription, or "STATs." There are six members of the STATs family. Stat1 and Stat3 are present in many cell types, as is Stat2 (as response to IFN-alpha is widespread). Stat4 is more restricted and is not in many cell types though it has been found in T helper class I, cells after treatment with IL-12. Stat5

was originally called mammary growth factor, but has been found at higher concentrations in other cells including myeloid cells. It can be activated in tissue culture cells by many cytokines.

[1113] The STATs are activated to translocate from the cytoplasm to the nucleus upon tyrosine phosphorylation by a set of kinases known as the Janus Kinase ("Jaks") family. Jaks represent a distinct family of soluble tyrosine kinases and include Tyk2, Jak1, Jak2, and Jak3. These kinases display significant sequence similarity and are generally catalytically inactive in resting cells.

[1114] The Jaks are activated by a wide range of receptors summarized in the Table below. (Adapted from review by Schidler and Darnell, *Ann. Rev. Biochem.* 64:621-51 (1995)). A cytokine receptor family, capable of activating Jaks, is divided into two groups: (a) Class 1 includes receptors for IL-2, IL-3, IL-4, IL-6, IL-7, IL-9, IL-11, IL-12, IL-15, Epo, PRL, GH, G-CSF, GM-CSF, LIF, CNTF, and thrombopoietin; and (b) Class 2 includes IFN- α , IFN- γ , and IL-10. The Class 1 receptors share a conserved cysteine motif (a set of four conserved cysteines and one tryptophan) and a WSXWS motif (a membrane proximal region encoding Trp-Ser-Xaa-Trp-Ser (SEQ ID NO: 2)).

[1115] Thus, on binding of a ligand to a receptor, Jaks are activated, which in turn activate STATs, which then translocate and bind to GAS elements. This entire process is encompassed in the Jaks-STATs signal transduction pathway.

[1116] Therefore, activation of the Jaks-STATs pathway, reflected by the binding of the GAS or the ISRE element, can be used to indicate proteins involved in the proliferation and differentiation of cells. For example, growth factors and cytokines are known to activate the Jaks-STATs pathway (See Table below). Thus, by using GAS elements linked to reporter molecules, activators of the Jaks-STATs pathway can be identified.

<u>Ligand</u>	<u>tyk2</u>	<u>JAKs</u> <u>Jak1</u>	<u>Jak2</u>	<u>Jak3</u>	<u>STATS</u>	<u>GAS(elements) or ISRE</u>
<u>IFN family</u>						
IFN-a/B	+	+	-	-	1,2,3	ISRE
IFN-g	+	+	-	-	1	GAS (IRF1>Lys6>IFP)
IL-10	+	?	?	-	1,3	
<u>gp130 family</u>						
IL-6 (Pleiotropic)	+	+	+	?	1,3	GAS (IRF1>Lys6>IFP)
IL-11(Pleiotropic)	?	+	?	?	1,3	
OnM(Pleiotropic)	?	+	+	?	1,3	
LIF(Pleiotropic)	?	+	+	?	1,3	
CNTF(Pleiotropic)	-/+	+	+	?	1,3	
G-CSF(Pleiotropic)	?	+	?	?	1,3	
IL-12(Pleiotropic)	+	-	+	+	1,3	
<u>g-C family</u>						
IL-2 (lymphocytes)	-	+	-	+	1,3,5	GAS
IL-4 (lymph/myeloid)	-	+	-	+	6	GAS (IRF1 = IFP
>>Ly6)(IgH)						
IL-7 (lymphocytes)	-	+	-	+	5	GAS
IL-9 (lymphocytes)	-	+	-	+	5	GAS
IL-13 (lymphocyte)	-	+	?	?	6	GAS
IL-15	?	+	?	+	5	GAS
<u>gp140 family</u>						
IL-3 (myeloid)	-	-	+	-	5	GAS (IRF1>IFP>>Ly6)
IL-5 (myeloid)	-	-	+	-	5	GAS
GM-CSF (myeloid)	-	-	+	-	5	GAS
<u>Growth hormone family</u>						
GH	?	-	+	-	5	
PRL	?	+/-	+	-	1,3,5	
EPO	?	-	+	-	5	GAS(B-
CAS>IRF1=IFP>>Ly6)						
<u>Receptor Tyrosine Kinases</u>						
EGF	?	+	+	-	1,3	GAS (IRF1)
PDGF	?	+	+	-	1,3	
CSF-1	?	+	+	-	1,3	GAS (not IRF1)

[1117] To construct a synthetic GAS containing promoter element, which is used in the Biological Assays described in Examples 32-33, a PCR based strategy is employed to generate a GAS-SV40 promoter sequence. The 5' primer contains four tandem copies of the GAS binding site found in the IRF1 promoter and previously demonstrated to bind STATs upon induction with a range of cytokines (Rothman et al., *Immunity* 1:457-468 (1994).), although other GAS or ISRE elements can be used instead. The 5' primer also contains 18bp of sequence complementary to the SV40 early promoter sequence and is flanked with an XhoI site. The sequence of the 5' primer is:

5':GCGCCTCGAGATTTCCCCGAAATCTAGATTTCCCCGAAATGATTTCCCCG
AAATGATTTCCCCGAAATATCTGCCATCTCAATTAG:3' (SEQ ID NO: 3)

[1118] The downstream primer is complementary to the SV40 promoter and is flanked with a Hind III site: 5':GCGGCAAGCTTTTTGCAAAGCCTAGGC:3' (SEQ ID NO: 4)

[1119] PCR amplification is performed using the SV40 promoter template present in the B-gal:promoter plasmid obtained from Clontech. The resulting PCR fragment is digested with XhoI/Hind III and subcloned into BLSK2-. (Stratagene.) Sequencing with forward and reverse primers confirms that the insert contains the following sequence:

5':CTCGAGATTTCCCCGAAATCTAGATTTCCCCGAAATGATTTCCCCGAAAT
GATTTCCCCGAAATATCTGCCATCTCAATTAGTCAGCAACCATAGTCCCGC
CCCTAACTCCGCCCATCCCGCCCCTAACTCCGCCCAGTTCCGCCCATTCTCC
GCCCCATGGCTGACTAATTTTTTTTATTTATGCAGAGGCCGAGGCCGCCTC
GGCCTCTGAGCTATTCCAGAAGTAGTGAGGAGGCTTTTTTGGAGGCCTAGG
CTTTTGCAAAAAGCTT:3' (SEQ ID NO: 5)

[1120] With this GAS promoter element linked to the SV40 promoter, a GAS:SEAP2 reporter construct is next engineered. Here, the reporter molecule is a secreted alkaline phosphatase, or "SEAP." Clearly, however, any reporter molecule can be instead of SEAP, in this or in any of the other Examples. Well known reporter molecules that can be used instead of SEAP include chloramphenicol acetyltransferase (CAT), luciferase, alkaline phosphatase, B-galactosidase, green fluorescent protein (GFP), or any protein detectable by an antibody.

[1121] The above sequence confirmed synthetic GAS-SV40 promoter element is subcloned into the pSEAP-Promoter vector obtained from Clontech using HindIII and XhoI, effectively replacing the SV40 promoter with the amplified GAS:SV40 promoter

element, to create the GAS-SEAP vector. However, this vector does not contain a neomycin resistance gene, and therefore, is not preferred for mammalian expression systems.

[1122] Thus, in order to generate mammalian stable cell lines expressing the GAS-SEAP reporter, the GAS-SEAP cassette is removed from the GAS-SEAP vector using SalI and NotI, and inserted into a backbone vector containing the neomycin resistance gene, such as pGFP-1 (Clontech), using these restriction sites in the multiple cloning site, to create the GAS-SEAP/Neo vector. Once this vector is transfected into mammalian cells, this vector can then be used as a reporter molecule for GAS binding as described in Examples 32-33.

[1123] Other constructs can be made using the above description and replacing GAS with a different promoter sequence. For example, construction of reporter molecules containing EGR and NF-KB promoter sequences are described in Examples 34 and 35. However, many other promoters can be substituted using the protocols described in these Examples. For instance, SRE, IL-2, NFAT, or Osteocalcin promoters can be substituted, alone or in combination (e.g., GAS/NF-KB/EGR, GAS/NF-KB, IL-2/NFAT, or NF-KB/GAS). Similarly, other cell lines can be used to test reporter construct activity, such as HELA (epithelial), HUVEC (endothelial), Reh (B-cell), Saos-2 (osteoblast), HUVAC (aortic), or Cardiomyocyte.

Example 32: High-Throughput Screening Assay for T-cell Activity.

[1124] The following protocol is used to assess T-cell activity by identifying factors, and determining whether supernate containing a polypeptide of the invention proliferates and/or differentiates T-cells. T-cell activity is assessed using the GAS/SEAP/Neo construct produced in Example 31. Thus, factors that increase SEAP activity indicate the ability to activate the Jaks-STATS signal transduction pathway. The T-cell used in this assay is Jurkat T-cells (ATCC Accession No. TIB-152), although Molt-3 cells (ATCC Accession No. CRL-1552) and Molt-4 cells (ATCC Accession No. CRL-1582) cells can also be used.

[1125] Jurkat T-cells are lymphoblastic CD4+ Th1 helper cells. In order to generate stable cell lines, approximately 2 million Jurkat cells are transfected with the GAS-

SEAP/neo vector using DMRIE-C (Life Technologies)(transfection procedure described below). The transfected cells are seeded to a density of approximately 20,000 cells per well and transfectants resistant to 1 mg/ml gentamicin selected. Resistant colonies are expanded and then tested for their response to increasing concentrations of interferon gamma. The dose response of a selected clone is demonstrated.

[1126] Specifically, the following protocol will yield sufficient cells for 75 wells containing 200 ul of cells. Thus, it is either scaled up, or performed in multiple to generate sufficient cells for multiple 96 well plates. Jurkat cells are maintained in RPMI + 10% serum with 1% Pen-Strep. Combine 2.5 mls of OPTI-MEM (Life Technologies) with 10 ug of plasmid DNA in a T25 flask. Add 2.5 ml OPTI-MEM containing 50 ul of DMRIE-C and incubate at room temperature for 15-45 mins.

[1127] During the incubation period, count cell concentration, spin down the required number of cells (10^7 per transfection), and resuspend in OPTI-MEM to a final concentration of 10^7 cells/ml. Then add 1ml of 1×10^7 cells in OPTI-MEM to T25 flask and incubate at 37 degree C for 6 hrs. After the incubation, add 10 ml of RPMI + 15% serum.

[1128] The Jurkat:GAS-SEAP stable reporter lines are maintained in RPMI + 10% serum, 1 mg/ml Gentamicin, and 1% Pen-Strep. These cells are treated with supernatants containing polypeptide of the present invention or polypeptide of the present invention induced polypeptides as produced by the protocol described in Example 30.

[1129] On the day of treatment with the supernatant, the cells should be washed and resuspended in fresh RPMI + 10% serum to a density of 500,000 cells per ml. The exact number of cells required will depend on the number of supernatants being screened. For one 96 well plate, approximately 10 million cells (for 10 plates, 100 million cells) are required.

[1130] Transfer the cells to a triangular reservoir boat, in order to dispense the cells into a 96 well dish, using a 12 channel pipette. Using a 12 channel pipette, transfer 200 ul of cells into each well (therefore adding 100,000 cells per well).

[1131] After all the plates have been seeded, 50 ul of the supernatants are transferred directly from the 96 well plate containing the supernatants into each well using a 12 channel pipette. In addition, a dose of exogenous interferon gamma (0.1, 1.0, 10 ng) is added to wells H9, H10, and H11 to serve as additional positive controls for the assay.

[1132] The 96 well dishes containing Jurkat cells treated with supernatants are placed in an incubator for 48 hrs (note: this time is variable between 48-72 hrs). 35 ul samples from each well are then transferred to an opaque 96 well plate using a 12 channel pipette. The opaque plates should be covered (using sellophene covers) and stored at -20 degree C until SEAP assays are performed according to Example 36. The plates containing the remaining treated cells are placed at 4 degree C and serve as a source of material for repeating the assay on a specific well if desired.

[1133] As a positive control, 100 Unit/ml interferon gamma can be used which is known to activate Jurkat T cells. Over 30 fold induction is typically observed in the positive control wells.

[1134] The above protocol may be used in the generation of both transient, as well as, stable transfected cells, which would be apparent to those of skill in the art.

Example 33: High-Throughput Screening Assay Identifying Myeloid Activity

[1135] The following protocol is used to assess myeloid activity of polypeptide of the present invention by determining whether polypeptide of the present invention proliferates and/or differentiates myeloid cells. Myeloid cell activity is assessed using the GAS/SEAP/Neo construct produced in Example 31. Thus, factors that increase SEAP activity indicate the ability to activate the Jaks-STATS signal transduction pathway. The myeloid cell used in this assay is U937, a pre-monocyte cell line, although TF-1, HL60, or KG1 can be used.

[1136] To transiently transfect U937 cells with the GAS/SEAP/Neo construct produced in Example 31, a DEAE-Dextran method (Kharbanda et. al., 1994, Cell Growth & Differentiation, 5:259-265) is used. First, harvest 2×10^7 U937 cells and wash with PBS. The U937 cells are usually grown in RPMI 1640 medium containing 10% heat-inactivated fetal bovine serum (FBS) supplemented with 100 units/ml penicillin and 100 mg/ml streptomycin.

[1137] Next, suspend the cells in 1 ml of 20 mM Tris-HCl (pH 7.4) buffer containing 0.5 mg/ml DEAE-Dextran, 8 ug GAS-SEAP2 plasmid DNA, 140 mM NaCl, 5 mM KCl,

375 μM $\text{Na}_2\text{HPO}_4 \cdot 7\text{H}_2\text{O}$, 1 mM MgCl_2 , and 675 μM CaCl_2 . Incubate at 37 degrees C for 45 min.

[1138] Wash the cells with RPMI 1640 medium containing 10% FBS and then resuspend in 10 ml complete medium and incubate at 37 degree C for 36 hr.

[1139] The GAS-SEAP/U937 stable cells are obtained by growing the cells in 400 $\mu\text{g}/\text{ml}$ G418. The G418-free medium is used for routine growth but every one to two months, the cells should be re-grown in 400 $\mu\text{g}/\text{ml}$ G418 for couple of passages.

[1140] These cells are tested by harvesting 1×10^8 cells (this is enough for ten 96-well plates assay) and wash with PBS. Suspend the cells in 200 ml above described growth medium, with a final density of 5×10^5 cells/ml. Plate 200 μl cells per well in the 96-well plate (or 1×10^5 cells/well).

[1141] Add 50 μl of the supernatant prepared by the protocol described in Example 30. Incubate at 37 degree C for 48 to 72 hr. As a positive control, 100 Unit/ml interferon gamma can be used which is known to activate U937 cells. Over 30 fold induction is typically observed in the positive control wells. SEAP assay the supernatant according to the protocol described in Example 36.

Example 34: High-Throughput Screening Assay Identifying Neuronal Activity.

[1142] When cells undergo differentiation and proliferation, a group of genes are activated through many different signal transduction pathways. One of these genes, EGR1 (early growth response gene 1), is induced in various tissues and cell types upon activation. The promoter of EGR1 is responsible for such induction. Using the EGR1 promoter linked to reporter molecules, activation of cells can be assessed by polypeptide of the present invention.

[1143] Particularly, the following protocol is used to assess neuronal activity in PC12 cell lines. PC12 cells (rat pheochromocytoma cells) are known to proliferate and/or differentiate by activation with a number of mitogens, such as TPA (tetradecanoyl phorbol acetate), NGF (nerve growth factor), and EGF (epidermal growth factor). The EGR1 gene expression is activated during this treatment. Thus, by stably transfecting PC12 cells with

a construct containing an EGR promoter linked to SEAP reporter, activation of PC12 cells by polypeptide of the present invention can be assessed.

[1144] The EGR/SEAP reporter construct can be assembled by the following protocol. The EGR-1 promoter sequence (-633 to +1)(Sakamoto K et al., Oncogene 6:867-871 (1991)) can be PCR amplified from human genomic DNA using the following primers:

5' GCGCTCGAGGGATGACAGCGATAGAACCCCGG-3' (SEQ ID NO: 6)

5' GCGAAGCTTCGCGACTCCCCGGATCCGCCTC-3' (SEQ ID NO: 7)

[1145] Using the GAS:SEAP/Neo vector produced in Example 31, EGR1 amplified product can then be inserted into this vector. Linearize the GAS:SEAP/Neo vector using restriction enzymes XhoI/HindIII, removing the GAS/SV40 stuffer. Restrict the EGR1 amplified product with these same enzymes. Ligate the vector and the EGR1 promoter.

[1146] To prepare 96 well-plates for cell culture, two mls of a coating solution (1:30 dilution of collagen type I (Upstate Biotech Inc. Cat#08-115) in 30% ethanol (filter sterilized)) is added per one 10 cm plate or 50 ml per well of the 96-well plate, and allowed to air dry for 2 hr.

[1147] PC12 cells are routinely grown in RPMI-1640 medium (Bio Whittaker) containing 10% horse serum (JRH BIOSCIENCES, Cat. # 12449-78P), 5% heat-inactivated fetal bovine serum (FBS) supplemented with 100 units/ml penicillin and 100 ug/ml streptomycin on a precoated 10 cm tissue culture dish. One to four split is done every three to four days. Cells are removed from the plates by scraping and resuspended with pipetting up and down for more than 15 times.

[1148] Transfect the EGR/SEAP/Neo construct into PC12 using the Lipofectamine protocol described in Example 30. EGR-SEAP/PC12 stable cells are obtained by growing the cells in 300 ug/ml G418. The G418-free medium is used for routine growth but every one to two months, the cells should be re-grown in 300 ug/ml G418 for couple of passages.

[1149] To assay for neuronal activity, a 10 cm plate with cells around 70 to 80% confluent is screened by removing the old medium. Wash the cells once with PBS (Phosphate buffered saline). Then starve the cells in low serum medium (RPMI-1640 containing 1% horse serum and 0.5% FBS with antibiotics) overnight.

[1150] The next morning, remove the medium and wash the cells with PBS. Scrape off the cells from the plate, suspend the cells well in 2 ml low serum medium. Count the cell number and add more low serum medium to reach final cell density as 5×10^5 cells/ml.

[1151] Add 200 μ l of the cell suspension to each well of 96-well plate (equivalent to 1×10^5 cells/well). Add 50 μ l supernatant produced by Example 30, 37 degree C for 48 to 72 hr. As a positive control, a growth factor known to activate PC12 cells through EGR can be used, such as 50 ng/ μ l of Neuronal Growth Factor (NGF). Over fifty-fold induction of SEAP is typically seen in the positive control wells. SEAP assay the supernatant according to Example 36.

Example 35: High-Throughput Screening Assay for T-cell Activity

[1152] NF-KB (Nuclear Factor KB) is a transcription factor activated by a wide variety of agents including the inflammatory cytokines IL-1 and TNF, CD30 and CD40, lymphotoxin-alpha and lymphotoxin-beta, by exposure to LPS or thrombin, and by expression of certain viral gene products. As a transcription factor, NF-KB regulates the expression of genes involved in immune cell activation, control of apoptosis (NF-KB appears to shield cells from apoptosis), B and T-cell development, anti-viral and antimicrobial responses, and multiple stress responses.

[1153] In non-stimulated conditions, NF-KB is retained in the cytoplasm with I-KB (Inhibitor KB). However, upon stimulation, I-KB is phosphorylated and degraded, causing NF-KB to shuttle to the nucleus, thereby activating transcription of target genes. Target genes activated by NF-KB include IL-2, IL-6, GM-CSF, ICAM-1 and class 1 MHC.

[1154] Due to its central role and ability to respond to a range of stimuli, reporter constructs utilizing the NF-KB promoter element are used to screen the supernatants produced in Example 30. Activators or inhibitors of NF-KB would be useful in treating, preventing, and/or diagnosing diseases. For example, inhibitors of NF-KB could be used to treat those diseases related to the acute or chronic activation of NF-KB, such as rheumatoid arthritis.

[1155] To construct a vector containing the NF-KB promoter element, a PCR based strategy is employed. The upstream primer contains four tandem copies of the NF-KB

binding site (GGGGACTTTCCC) (SEQ ID NO: 8), 18 bp of sequence complementary to the 5' end of the SV40 early promoter sequence, and is flanked with an XhoI site:

5':GCGGCCTCGAGGGGACTTTCCCGGGGACTTTCCGGGGACTTTCCGGGAC
TTTCCATCTGCCATCTCAATTAG:3' (SEQ ID NO: 9)

[1156] The downstream primer is complementary to the 3' end of the SV40 promoter and is flanked with a Hind III site:

5':GCGGCAAGCTTTTTGCAAAGCCTAGGC:3' (SEQ ID NO: 4)

[1157] PCR amplification is performed using the SV40 promoter template present in the pB-gal:promoter plasmid obtained from Clontech. The resulting PCR fragment is digested with XhoI and Hind III and subcloned into BLSK2-. (Stratagene) Sequencing with the T7 and T3 primers confirms the insert contains the following sequence:

5':CTCGAGGGGACTTTCCCGGGGACTTTCCGGGGACTTTCCGGGACTTTCC
ATCTGCCATCTCAATTAGTCAGCAACCATAGTCCCGCCCTAACTCCGCCC
ATCCCGCCCCCTAACTCCGCCCAGTTCCGCCCATTCTCCGCCCCATGGCTGA
CTAATTTTTTTTATTTATGCAGAGGCCGAGGCCGCCTCGGCCTCTGAGCTAT
TCCAGAAGTAGTGAGGAGGCTTTTTTGGAGGCCTAGGCTTTTGCAAAAAGC
TT:3' (SEQ ID NO: 10)

[1158] Next, replace the SV40 minimal promoter element present in the pSEAP2-promoter plasmid (Clontech) with this NF-KB/SV40 fragment using XhoI and HindIII. However, this vector does not contain a neomycin resistance gene, and therefore, is not preferred for mammalian expression systems.

[1159] In order to generate stable mammalian cell lines, the NF-KB/SV40/SEAP cassette is removed from the above NF-KB/SEAP vector using restriction enzymes SalI and NotI, and inserted into a vector containing neomycin resistance. Particularly, the NF-KB/SV40/SEAP cassette was inserted into pGFP-1 (Clontech), replacing the GFP gene, after restricting pGFP-1 with SalI and NotI.

[1160] Once NF-KB/SV40/SEAP/Neo vector is created, stable Jurkat T-cells are created and maintained according to the protocol described in Example 32. Similarly, the method for assaying supernatants with these stable Jurkat T-cells is also described in Example 32. As a positive control, exogenous TNF alpha (0.1, 1, 10 ng) is added to wells H9, H10, and H11, with a 5-10 fold activation typically observed.

Example 36: Assay for SEAP Activity

[1161] As a reporter molecule for the assays described in Examples 32-35, SEAP activity is assayed using the Tropix Phospho-light Kit (Cat. BP-400) according to the following general procedure. The Tropix Phospho-light Kit supplies the Dilution, Assay, and Reaction Buffers used below.

[1162] Prime a dispenser with the 2.5x Dilution Buffer and dispense 15 ul of 2.5x dilution buffer into Optiplates containing 35 ul of a supernatant. Seal the plates with a plastic sealer and incubate at 65 degree C for 30 min. Separate the Optiplates to avoid uneven heating.

[1163] Cool the samples to room temperature for 15 minutes. Empty the dispenser and prime with the Assay Buffer. Add 50 ml Assay Buffer and incubate at room temperature 5 min. Empty the dispenser and prime with the Reaction Buffer (see the Table below). Add 50 ul Reaction Buffer and incubate at room temperature for 20 minutes. Since the intensity of the chemiluminescent signal is time dependent, and it takes about 10 minutes to read 5 plates on a luminometer, thus one should treat 5 plates at each time and start the second set 10 minutes later.

[1164] Read the relative light unit in the luminometer. Set H12 as blank, and print the results. An increase in chemiluminescence indicates reporter activity.

Reaction Buffer Formulation:

# of plates	Rxn buffer diluent (ml)	CSPD (ml)
10	60	3
11	65	3.25
12	70	3.5
13	75	3.75
14	80	4
15	85	4.25
16	90	4.5
17	95	4.75
18	100	5
19	105	5.25
20	110	5.5
21	115	5.75

22	120	6
23	125	6.25
24	130	6.5
25	135	6.75
26	140	7
27	145	7.25
28	150	7.5
29	155	7.75
30	160	8
31	165	8.25
32	170	8.5
33	175	8.75
34	180	9
35	185	9.25
36	190	9.5
37	195	9.75
38	200	10
39	205	10.25
40	210	10.5
41	215	10.75
42	220	11
43	225	11.25
44	230	11.5
45	235	11.75
46	240	12
47	245	12.25
48	250	12.5
49	255	12.75
50	260	13

***Example 37: High-Throughput Screening Assay Identifying Changes in
Small Molecule Concentration and Membrane Permeability***

[1165] Binding of a ligand to a receptor is known to alter intracellular levels of small molecules, such as calcium, potassium, sodium, and pH, as well as alter membrane potential. These alterations can be measured in an assay to identify supernatants which

bind to receptors of a particular cell. Although the following protocol describes an assay for calcium, this protocol can easily be modified to detect changes in potassium, sodium, pH, membrane potential, or any other small molecule which is detectable by a fluorescent probe.

[1166] The following assay uses Fluorometric Imaging Plate Reader ("FLIPR") to measure changes in fluorescent molecules (Molecular Probes) that bind small molecules. Clearly, any fluorescent molecule detecting a small molecule can be used instead of the calcium fluorescent molecule, fluo-4 (Molecular Probes, Inc.; catalog no. F-14202), used here.

[1167] For adherent cells, seed the cells at 10,000 -20,000 cells/well in a Co-star black 96-well plate with clear bottom. The plate is incubated in a CO₂ incubator for 20 hours. The adherent cells are washed two times in Biotek washer with 200 ul of HBSS (Hank's Balanced Salt Solution) leaving 100 ul of buffer after the final wash.

[1168] A stock solution of 1 mg/ml fluo-4 is made in 10% pluronic acid DMSO. To load the cells with fluo-4, 50 ul of 12 ug/ml fluo-4 is added to each well. The plate is incubated at 37 degrees C in a CO₂ incubator for 60 min. The plate is washed four times in the Biotek washer with HBSS leaving 100 ul of buffer.

[1169] For non-adherent cells, the cells are spun down from culture media. Cells are re-suspended to 2.5×10^6 cells/ml with HBSS in a 50-ml conical tube. 4 ul of 1 mg/ml fluo-4 solution in 10% pluronic acid DMSO is added to each ml of cell suspension. The tube is then placed in a 37 degrees C water bath for 30-60 min. The cells are washed twice with HBSS, resuspended to 1×10^6 cells/ml, and dispensed into a microplate, 100 ul/well. The plate is centrifuged at 1000 rpm for 5 min. The plate is then washed once in Denley Cell Wash with 200 ul, followed by an aspiration step to 100 ul final volume.

[1170] For a non-cell based assay, each well contains a fluorescent molecule, such as fluo-4. The supernatant is added to the well, and a change in fluorescence is detected.

[1171] To measure the fluorescence of intracellular calcium, the FLIPR is set for the following parameters: (1) System gain is 300-800 mW; (2) Exposure time is 0.4 second; (3) Camera F/stop is F/2; (4) Excitation is 488 nm; (5) Emission is 530 nm; and (6) Sample addition is 50 ul. Increased emission at 530 nm indicates an extracellular signaling event caused by the a molecule, either polypeptide of the present invention or a

molecule induced by polypeptide of the present invention, which has resulted in an increase in the intracellular Ca^{++} concentration.

Example 38: High-Throughput Screening Assay Identifying Tyrosine Kinase Activity

[1172] The Protein Tyrosine Kinases (PTK) represent a diverse group of transmembrane and cytoplasmic kinases. Within the Receptor Protein Tyrosine Kinase (RPTK) group are receptors for a range of mitogenic and metabolic growth factors including the PDGF, FGF, EGF, NGF, HGF and Insulin receptor subfamilies. In addition there are a large family of RPTKs for which the corresponding ligand is unknown. Ligands for RPTKs include mainly secreted small proteins, but also membrane-bound and extracellular matrix proteins.

[1173] Activation of RPTK by ligands involves ligand-mediated receptor dimerization, resulting in transphosphorylation of the receptor subunits and activation of the cytoplasmic tyrosine kinases. The cytoplasmic tyrosine kinases include receptor associated tyrosine kinases of the src-family (e.g., src, yes, lck, lyn, fyn) and non-receptor linked and cytosolic protein tyrosine kinases, such as the Jak family, members of which mediate signal transduction triggered by the cytokine superfamily of receptors (e.g., the Interleukins, Interferons, GM-CSF, and Leptin).

[1174] Because of the wide range of known factors capable of stimulating tyrosine kinase activity, identifying whether polypeptide of the present invention or a molecule induced by polypeptide of the present invention is capable of activating tyrosine kinase signal transduction pathways is of interest. Therefore, the following protocol is designed to identify such molecules capable of activating the tyrosine kinase signal transduction pathways.

[1175] Seed target cells (e.g., primary keratinocytes) at a density of approximately 25,000 cells per well in a 96 well Loprodyne Silent Screen Plates purchased from Nalge Nunc (Naperville, IL). The plates are sterilized with two 30 minute rinses with 100% ethanol, rinsed with water and dried overnight. Some plates are coated for 2 hr with 100 ml of cell culture grade type I collagen (50 mg/ml), gelatin (2%) or polylysine (50 mg/ml), all of which can be purchased from Sigma Chemicals (St. Louis, MO) or 10%

Matrigel purchased from Becton Dickinson (Bedford,MA), or calf serum, rinsed with PBS and stored at 4 degree C. Cell growth on these plates is assayed by seeding 5,000 cells/well in growth medium and indirect quantitation of cell number through use of alamarBlue as described by the manufacturer Alamar Biosciences, Inc. (Sacramento, CA) after 48 hr. Falcon plate covers #3071 from Becton Dickinson (Bedford,MA) are used to cover the Loprodyne Silent Screen Plates. Falcon Microtest III cell culture plates can also be used in some proliferation experiments.

[1176] To prepare extracts, A431 cells are seeded onto the nylon membranes of Loprodyne plates (20,000/200ml/well) and cultured overnight in complete medium. Cells are quiesced by incubation in serum-free basal medium for 24 hr. After 5-20 minutes treatment with EGF (60ng/ml) or 50 ul of the supernatant produced in Example 30, the medium was removed and 100 ml of extraction buffer ((20 mM HEPES pH 7.5, 0.15 M NaCl, 1% Triton X-100, 0.1% SDS, 2 mM Na₃VO₄, 2 mM Na₄P₂O₇ and a cocktail of protease inhibitors (# 1836170) obtained from Boehringer Mannheim (Indianapolis, IN)) is added to each well and the plate is shaken on a rotating shaker for 5 minutes at 4°C. The plate is then placed in a vacuum transfer manifold and the extract filtered through the 0.45 mm membrane bottoms of each well using house vacuum. Extracts are collected in a 96-well catch/assay plate in the bottom of the vacuum manifold and immediately placed on ice. To obtain extracts clarified by centrifugation, the content of each well, after detergent solubilization for 5 minutes, is removed and centrifuged for 15 minutes at 4 degree C at 16,000 x g.

[1177] Test the filtered extracts for levels of tyrosine kinase activity. Although many methods of detecting tyrosine kinase activity are known, one method is described here.

[1178] Generally, the tyrosine kinase activity of a supernatant is evaluated by determining its ability to phosphorylate a tyrosine residue on a specific substrate (a biotinylated peptide). Biotinylated peptides that can be used for this purpose include PSK1 (corresponding to amino acids 6-20 of the cell division kinase cdc2-p34) and PSK2 (corresponding to amino acids 1-17 of gastrin). Both peptides are substrates for a range of tyrosine kinases and are available from Boehringer Mannheim.

[1179] The tyrosine kinase reaction is set up by adding the following components in order. First, add 10ul of 5uM Biotinylated Peptide, then 10ul ATP/Mg₂₊ (5mM ATP/50mM MgCl₂), then 10ul of 5x Assay Buffer (40mM imidazole hydrochloride,

pH7.3, 40 mM beta-glycerophosphate, 1mM EGTA, 100mM MgCl₂, 5 mM MnCl₂, 0.5 mg/ml BSA), then 5ul of Sodium Vanadate(1mM), and then 5ul of water. Mix the components gently and preincubate the reaction mix at 30 degree C for 2 min. Initiate the reaction by adding 10ul of the control enzyme or the filtered supernatant.

[1180] The tyrosine kinase assay reaction is then terminated by adding 10 ul of 120mM EDTA and place the reactions on ice.

[1181] Tyrosine kinase activity is determined by transferring 50 ul aliquot of reaction mixture to a microtiter plate (MTP) module and incubating at 37 degree C for 20 min. This allows the streptavidin coated 96 well plate to associate with the biotinylated peptide. Wash the MTP module with 300ul/well of PBS four times. Next add 75 ul of anti-phosphotyrosine antibody conjugated to horse radish peroxidase(anti-P-Tyr-POD(0.5u/ml)) to each well and incubate at 37 degree C for one hour. Wash the well as above.

[1182] Next add 100ul of peroxidase substrate solution (Boehringer Mannheim) and incubate at room temperature for at least 5 mins (up to 30 min). Measure the absorbance of the sample at 405 nm by using ELISA reader. The level of bound peroxidase activity is quantitated using an ELISA reader and reflects the level of tyrosine kinase activity.

Example 39: High-Throughput Screening Assay Identifying

Phosphorylation Activity

[1183] As a potential alternative and/or complement to the assay of protein tyrosine kinase activity described in Example 38, an assay which detects activation (phosphorylation) of major intracellular signal transduction intermediates can also be used. For example, as described below one particular assay can detect tyrosine phosphorylation of the Erk-1 and Erk-2 kinases. However, phosphorylation of other molecules, such as Raf, JNK, p38 MAP, Map kinase kinase (MEK), MEK kinase, Src, Muscle specific kinase (MuSK), IRAK, Tec, and Janus, as well as any other phosphoserine, phosphotyrosine, or phosphothreonine molecule, can be detected by substituting these molecules for Erk-1 or Erk-2 in the following assay.

[1184] Specifically, assay plates are made by coating the wells of a 96-well ELISA plate with 0.1ml of protein G (1ug/ml) for 2 hr at room temp, (RT). The plates are then rinsed with PBS and blocked with 3% BSA/PBS for 1 hr at RT. The protein G plates are

then treated with 2 commercial monoclonal antibodies (100ng/well) against Erk-1 and Erk-2 (1 hr at RT) (Santa Cruz Biotechnology). (To detect other molecules, this step can easily be modified by substituting a monoclonal antibody detecting any of the above described molecules.) After 3-5 rinses with PBS, the plates are stored at 4 degree C until use.

[1185] A431 cells are seeded at 20,000/well in a 96-well Loprodyne filterplate and cultured overnight in growth medium. The cells are then starved for 48 hr in basal medium (DMEM) and then treated with EGF (6ng/well) or 50 ul of the supernatants obtained in Example 30 for 5-20 minutes. The cells are then solubilized and extracts filtered directly into the assay plate.

[1186] After incubation with the extract for 1 hr at RT, the wells are again rinsed. As a positive control, a commercial preparation of MAP kinase (10ng/well) is used in place of A431 extract. Plates are then treated with a commercial polyclonal (rabbit) antibody (1ug/ml) which specifically recognizes the phosphorylated epitope of the Erk-1 and Erk-2 kinases (1 hr at RT). This antibody is biotinylated by standard procedures. The bound polyclonal antibody is then quantitated by successive incubations with Europium-streptavidin and Europium fluorescence enhancing reagent in the Wallac DELFIA instrument (time-resolved fluorescence). An increased fluorescent signal over background indicates a phosphorylation by polypeptide of the present invention or a molecule induced by polypeptide of the present invention.

Example 40: Assay for the Stimulation of Bone Marrow CD34+ Cell Proliferation

[1187] This assay is based on the ability of human CD34+ to proliferate in the presence of hematopoietic growth factors and evaluates the ability of isolated polypeptides expressed in mammalian cells to stimulate proliferation of CD34+ cells.

[1188] It has been previously shown that most mature precursors will respond to only a single signal. More immature precursors require at least two signals to respond. Therefore, to test the effect of polypeptides on hematopoietic activity of a wide range of progenitor cells, the assay contains a given polypeptide in the presence or absence of other hematopoietic growth factors. Isolated cells are cultured for 5 days in the presence of

Stem Cell Factor (SCF) in combination with tested sample. SCF alone has a very limited effect on the proliferation of bone marrow (BM) cells, acting in such conditions only as a "survival" factor. However, combined with any factor exhibiting stimulatory effect on these cells (e.g., IL-3), SCF will cause a synergistic effect. Therefore, if the tested polypeptide has a stimulatory effect on hematopoietic progenitors, such activity can be easily detected. Since normal BM cells have a low level of cycling cells, it is likely that any inhibitory effect of a given polypeptide, or agonists or antagonists thereof, might not be detected. Accordingly, assays for an inhibitory effect on progenitors is preferably tested in cells that are first subjected to *in vitro* stimulation with SCF+IL-3, and then contacted with the compound that is being evaluated for inhibition of such induced proliferation.

[1189] Briefly, CD34+ cells are isolated using methods known in the art. The cells are thawed and resuspended in medium (QBSF 60 serum-free medium with 1% L-glutamine (500ml) Quality Biological, Inc., Gaithersburg, MD Cat# 160-204-101). After several gentle centrifugation steps at 200 x g, cells are allowed to rest for one hour. The cell count is adjusted to 2.5×10^5 cells/ml. During this time, 100 μ l of sterile water is added to the peripheral wells of a 96-well plate. The cytokines that can be tested with a given polypeptide in this assay is rhSCF (R&D Systems, Minneapolis, MN, Cat# 255-SC) at 50 ng/ml alone and in combination with rhSCF and rhIL-3 (R&D Systems, Minneapolis, MN, Cat# 203-ML) at 30 ng/ml. After one hour, 10 μ l of prepared cytokines, 50 μ l of the supernatants prepared in Example 30 (supernatants at 1:2 dilution = 50 μ l) and 20 μ l of diluted cells are added to the media which is already present in the wells to allow for a final total volume of 100 μ l. The plates are then placed in a 37°C/5% CO₂ incubator for five days.

[1190] Eighteen hours before the assay is harvested, 0.5 μ Ci/well of [3H] Thymidine is added in a 10 μ l volume to each well to determine the proliferation rate. The experiment is terminated by harvesting the cells from each 96-well plate to a filtermat using the Tomtec Harvester 96. After harvesting, the filtermats are dried, trimmed and placed into OmniFilter assemblies consisting of one OmniFilter plate and one OmniFilter Tray. 60 μ l Microscint is added to each well and the plate sealed with TopSeal-A press-on sealing film. A bar code 15 sticker is affixed to the first plate for counting. The sealed plates are then loaded and the level of radioactivity determined via the Packard Top Count and the

printed data collected for analysis. The level of radioactivity reflects the amount of cell proliferation.

[1191] The studies described in this example test the activity of a given polypeptide to stimulate bone marrow CD34+ cell proliferation. One skilled in the art could easily modify the exemplified studies to test the activity of polynucleotides (e.g., gene therapy), antibodies, agonists, and/or antagonists and fragments and variants thereof. As a nonlimiting example, potential antagonists tested in this assay would be expected to inhibit cell proliferation in the presence of cytokines and/or to increase the inhibition of cell proliferation in the presence of cytokines and a given polypeptide. In contrast, potential agonists tested in this assay would be expected to enhance cell proliferation and/or to decrease the inhibition of cell proliferation in the presence of cytokines and a given polypeptide.

[1192] The ability of a gene to stimulate the proliferation of bone marrow CD34+ cells indicates that polynucleotides and polypeptides corresponding to the gene are useful for the diagnosis and treatment of disorders affecting the immune system and hematopoiesis. Representative uses are described in the "Immune Activity" and "Infectious Disease" sections above, and elsewhere herein.

[1193] ***Example 41: Assay for Extracellular Matrix Enhanced Cell Response (EMECCR)***

[1194] The objective of the Extracellular Matrix Enhanced Cell Response (EMECCR) assay is to identify gene products (e.g., isolated polypeptides) that act on the hematopoietic stem cells in the context of the extracellular matrix (ECM) induced signal.

[1195] Cells respond to the regulatory factors in the context of signal(s) received from the surrounding microenvironment. For example, fibroblasts, and endothelial and epithelial stem cells fail to replicate in the absence of signals from the ECM. Hematopoietic stem cells can undergo self-renewal in the bone marrow, but not in *in vitro* suspension culture. The ability of stem cells to undergo self-renewal *in vitro* is dependent upon their interaction with the stromal cells and the ECM protein fibronectin (fn). Adhesion of cells to fn is mediated by the $\alpha_5\beta_1$ and $\alpha_4\beta_1$ integrin receptors, which are expressed by human and mouse hematopoietic stem cells. The factor(s) which integrate

with the ECM environment and are responsible for stimulating stem cell self-renewal have not yet been identified. Discovery of such factors should be of great interest in gene therapy and bone marrow transplant applications

[1196] Briefly, polystyrene, non tissue culture treated, 96-well plates are coated with fragment at a coating concentration of $0.2 \mu\text{g}/\text{cm}^2$. Mouse bone marrow cells are plated (1,000 cells/well) in 0.2 ml of serum-free medium. Cells cultured in the presence of IL-3 (5 ng/ml) + SCF (50 ng/ml) would serve as the positive control, conditions under which little self-renewal but pronounced differentiation of the stem cells is to be expected. Gene products of the invention (e.g., including, but not limited to, polynucleotides and polypeptides of the present invention, and supernatants produced in Example 30), are tested with appropriate negative controls in the presence and absence of SCF(5.0 ng/ml), where test factor supernatants represent 10% of the total assay volume. The plated cells are then allowed to grow by incubating in a low oxygen environment (5% CO_2 , 7% O_2 , and 88% N_2) tissue culture incubator for 7 days. The number of proliferating cells within the wells is then quantitated by measuring thymidine incorporation into cellular DNA. Verification of the positive hits in the assay will require phenotypic characterization of the cells, which can be accomplished by scaling up of the culture system and using appropriate antibody reagents against cell surface antigens and FACSscan.

[1197] One skilled in the art could easily modify the exemplified studies to test the activity of polynucleotides (e.g., gene therapy), antibodies, agonists, and/or antagonists and fragments and variants thereof.

[1198] If a particular polypeptide of the present invention is found to be a stimulator of hematopoietic progenitors, polynucleotides and polypeptides corresponding to the gene encoding said polypeptide may be useful for the diagnosis and treatment of disorders affecting the immune system and hematopoiesis. Representative uses are described in the "Immune Activity" and "Infectious Disease" sections above, and elsewhere herein. The gene product may also be useful in the expansion of stem cells and committed progenitors of various blood lineages, and in the differentiation and/or proliferation of various cell types.

[1199] Additionally, the polynucleotides and/or polypeptides of the gene of interest and/or agonists and/or antagonists thereof, may also be employed to inhibit the proliferation and differentiation of hematopoietic cells and therefore may be employed to

protect bone marrow stem cells from chemotherapeutic agents during chemotherapy. This antiproliferative effect may allow administration of higher doses of chemotherapeutic agents and, therefore, more effective chemotherapeutic treatment.

[1200] Moreover, polynucleotides and polypeptides corresponding to the gene of interest may also be useful for the treatment and diagnosis of hematopoietic related disorders such as, for example, anemia, pancytopenia, leukopenia, thrombocytopenia or leukemia since stromal cells are important in the production of cells of hematopoietic lineages. The uses include bone marrow cell ex-vivo culture, bone marrow transplantation, bone marrow reconstitution, radiotherapy or chemotherapy of neoplasia.

Example 42: Human Dermal Fibroblast and Aortic Smooth Muscle Cell Proliferation

[1201] The polypeptide of interest is added to cultures of normal human dermal fibroblasts (NHDF) and human aortic smooth muscle cells (AoSMC) and two co-assays are performed with each sample. The first assay examines the effect of the polypeptide of interest on the proliferation of normal human dermal fibroblasts (NHDF) or aortic smooth muscle cells (AoSMC). Aberrant growth of fibroblasts or smooth muscle cells is a part of several pathological processes, including fibrosis, and restenosis. The second assay examines IL6 production by both NHDF and SMC. IL6 production is an indication of functional activation. Activated cells will have increased production of a number of cytokines and other factors, which can result in a proinflammatory or immunomodulatory outcome. Assays are run with and without co-TNF α stimulation, in order to check for costimulatory or inhibitory activity.

[1202] Briefly, on day 1, 96-well black plates are set up with 1000 cells/well (NHDF) or 2000 cells/well (AoSMC) in 100 μ l culture media. NHDF culture media contains: Clonetics FB basal media, 1mg/ml hFGF, 5mg/ml insulin, 50mg/ml gentamycin, 2%FBS, while AoSMC culture media contains Clonetics SM basal media, 0.5 μ g/ml hEGF, 5mg/ml insulin, 1 μ g/ml hFGF, 50mg/ml gentamycin, 50 μ g/ml Amphotericin B, 5%FBS. After incubation at 37°C for at least 4-5 hours culture media is aspirated and replaced with growth arrest media. Growth arrest media for NHDF contains fibroblast basal media, 50mg/ml gentamycin, 2% FBS, while growth arrest media for AoSMC contains SM basal

media, 50mg/ml gentamycin, 50 μ g/ml Amphotericin B, 0.4% FBS. Incubate at 37 °C until day 2.

[1203] On day 2, serial dilutions and templates of the polypeptide of interest are designed such that they always include media controls and known-protein controls. For both stimulation and inhibition experiments, proteins are diluted in growth arrest media. For inhibition experiments, TNF α is added to a final concentration of 2ng/ml (NHDF) or 5ng/ml (AoSMC). Add 1/3 vol media containing controls or polypeptides of the present invention and incubate at 37 degrees C/5% CO₂ until day 5.

[1204] Transfer 60 μ l from each well to another labeled 96-well plate, cover with a plate-sealer, and store at 4 degrees C until Day 6 (for IL6 ELISA). To the remaining 100 μ l in the cell culture plate, aseptically add Alamar Blue in an amount equal to 10% of the culture volume (10 μ l). Return plates to incubator for 3 to 4 hours. Then measure fluorescence with excitation at 530nm and emission at 590nm using the CytoFluor. This yields the growth stimulation/inhibition data.

[1205] On day 5, the IL6 ELISA is performed by coating a 96 well plate with 50-100 μ l/well of Anti-Human IL6 Monoclonal antibody diluted in PBS, pH 7.4, incubate ON at room temperature.

[1206] On day 6, empty the plates into the sink and blot on paper towels. Prepare Assay Buffer containing PBS with 4% BSA. Block the plates with 200 μ l/well of Pierce Super Block blocking buffer in PBS for 1-2 hr and then wash plates with wash buffer (PBS, 0.05% Tween-20). Blot plates on paper towels. Then add 50 μ l/well of diluted Anti-Human IL-6 Monoclonal, Biotin-labeled antibody at 0.50 mg/ml. Make dilutions of IL-6 stock in media (30, 10, 3, 1, 0.3, 0 ng/ml). Add duplicate samples to top row of plate. Cover the plates and incubate for 2 hours at RT on shaker.

[1207] Plates are washed with wash buffer and blotted on paper towels. Dilute EU-labeled Streptavidin 1:1000 in Assay buffer, and add 100 μ l/well. Cover the plate and incubate 1 h at RT. Plates are again washed with wash buffer and blotted on paper towels.

[1208] Add 100 μ l/well of Enhancement Solution. Shake for 5 minutes. Read the plate on the Wallac DELFIA Fluorometer. Readings from triplicate samples in each assay were tabulated and averaged.

[1209] A positive result in this assay suggests AoSMC cell proliferation and that the polypeptide of the present invention may be involved in dermal fibroblast proliferation

and/or smooth muscle cell proliferation. A positive result also suggests many potential uses of polypeptides, polynucleotides, agonists and/or antagonists of the polynucleotide/polypeptide of the present invention which gives a positive result. For example, inflammation and immune responses, wound healing, and angiogenesis, as detailed throughout this specification. Particularly, polypeptides of the present invention and polynucleotides of the present invention may be used in wound healing and dermal regeneration, as well as the promotion of vasculogenesis, both of the blood vessels and lymphatics. The growth of vessels can be used in the treatment of, for example, cardiovascular diseases. Additionally, antagonists of polypeptides and polynucleotides of the invention may be useful in treating diseases, disorders, and/or conditions which involve angiogenesis by acting as an anti-vascular agent (e.g., anti-angiogenesis). These diseases, disorders, and/or conditions are known in the art and/or are described herein, such as, for example, malignancies, solid tumors, benign tumors, for example hemangiomas, acoustic neuromas, neurofibromas, trachomas, and pyogenic granulomas; arteriosclerotic plaques; ocular angiogenic diseases, for example, diabetic retinopathy, retinopathy of prematurity, macular degeneration, corneal graft rejection, neovascular glaucoma, retrolental fibroplasia, rubeosis, retinoblastoma, uveitis and Pterygia (abnormal blood vessel growth) of the eye; rheumatoid arthritis; psoriasis; delayed wound healing; endometriosis; vasculogenesis; granulations; hypertrophic scars (keloids); nonunion fractures; scleroderma; trachoma; vascular adhesions; myocardial angiogenesis; coronary collaterals; cerebral collaterals; arteriovenous malformations; ischemic limb angiogenesis; Osler-Webber Syndrome; plaque neovascularization; telangiectasia; hemophilic joints; angiofibroma; fibromuscular dysplasia; wound granulation; Crohn's disease; and atherosclerosis. Moreover, antagonists of polypeptides and polynucleotides of the invention may be useful in treating anti-hyperproliferative diseases and/or anti-inflammatory known in the art and/or described herein.

[1210] One skilled in the art could easily modify the exemplified studies to test the activity of polynucleotides (e.g., gene therapy), antibodies, agonists, and/or antagonists and fragments and variants thereof.

***Example 43: Cellular Adhesion Molecule (CAM) Expression on
Endothelial Cells***

[1211] The recruitment of lymphocytes to areas of inflammation and angiogenesis involves specific receptor-ligand interactions between cell surface adhesion molecules (CAMs) on lymphocytes and the vascular endothelium. The adhesion process, in both normal and pathological settings, follows a multi-step cascade that involves intercellular adhesion molecule-1 (ICAM-1), vascular cell adhesion molecule-1 (VCAM-1), and endothelial leukocyte adhesion molecule-1 (E-selectin) expression on endothelial cells (EC). The expression of these molecules and others on the vascular endothelium determines the efficiency with which leukocytes may adhere to the local vasculature and extravasate into the local tissue during the development of an inflammatory response. The local concentration of cytokines and growth factor participate in the modulation of the expression of these CAMs.

[1212] Briefly, endothelial cells (e.g., Human Umbilical Vein Endothelial cells (HUVECs)) are grown in a standard 96 well plate to confluence, growth medium is removed from the cells and replaced with 100 μ l of 199 Medium (10% fetal bovine serum (FBS)). Samples for testing and positive or negative controls are added to the plate in triplicate (in 10 μ l volumes). Plates are then incubated at 37°C for either 5 h (selectin and integrin expression) or 24 h (integrin expression only). Plates are aspirated to remove medium and 100 μ l of 0.1% paraformaldehyde-PBS(with Ca++ and Mg++) is added to each well. Plates are held at 4°C for 30 min. Fixative is removed from the wells and wells are washed 1X with PBS(+Ca,Mg) + 0.5% BSA and drained. 10 μ l of diluted primary antibody is added to the test and control wells. Anti-ICAM-1-Biotin, Anti-VCAM-1-Biotin and Anti-E-selectin-Biotin are used at a concentration of 10 μ g/ml (1:10 dilution of 0.1 mg/ml stock antibody). Cells are incubated at 37°C for 30 min. in a humidified environment. Wells are washed three times with PBS(+Ca,Mg) + 0.5% BSA. 20 μ l of diluted ExtrAvidin-Alkaline Phosphatase (1:5,000 dilution, referred to herein as the working dilution) are added to each well and incubated at 37°C for 30 min. Wells are washed three times with PBS(+Ca,Mg)+0.5% BSA. Dissolve 1 tablet of p-Nitrophenol Phosphate pNPP per 5 ml of glycine buffer (pH 10.4). 100 μ l of pNPP substrate in glycine buffer is added to each test well. Standard wells in triplicate are prepared from the

working dilution of the ExtraAvidin-Alkaline Phosphatase in glycine buffer: $1:5,000$ (10^0) $> 10^{-0.5} > 10^{-1} > 10^{-1.5}$. $5 \mu\text{l}$ of each dilution is added to triplicate wells and the resulting AP content in each well is 5.50 ng, 1.74 ng, 0.55 ng, 0.18 ng. $100 \mu\text{l}$ of pNNP reagent is then added to each of the standard wells. The plate is incubated at 37°C for 4h. A volume of $50 \mu\text{l}$ of 3M NaOH is added to all wells. The plate is read on a plate reader at 405 nm using the background subtraction option on blank wells filled with glycine buffer only. Additionally, the template is set up to indicate the concentration of AP-conjugate in each standard well [5.50 ng; 1.74 ng; 0.55 ng; 0.18 ng]. Results are indicated as amount of bound AP-conjugate in each sample.

Example 44: Alamar Blue Endothelial Cells Proliferation Assay

[1213] This assay may be used to quantitatively determine protein mediated inhibition of bFGF-induced proliferation of Bovine Lymphatic Endothelial Cells (LECs), Bovine Aortic Endothelial Cells (BAECs) or Human Microvascular Uterine Myometrial Cells (UTMECs). This assay incorporates a fluorometric growth indicator based on detection of metabolic activity. A standard Alamar Blue Proliferation Assay is prepared in EGM-2MV with 10 ng/ml of bFGF added as a source of endothelial cell stimulation. This assay may be used with a variety of endothelial cells with slight changes in growth medium and cell concentration. Dilutions of the protein batches to be tested are diluted as appropriate. Serum-free medium (GIBCO SFM) without bFGF is used as a non-stimulated control and Angiostatin or TSP-1 are included as a known inhibitory controls.

[1214] Briefly, LEC, BAECs or UTMECs are seeded in growth media at a density of 5000 to 2000 cells/well in a 96 well plate and placed at 37°C overnight. After the overnight incubation of the cells, the growth media is removed and replaced with GIBCO EC-SFM. The cells are treated with the appropriate dilutions of the protein of interest or control protein sample(s) (prepared in SFM) in triplicate wells with additional bFGF to a concentration of 10 ng/ml. Once the cells have been treated with the samples, the plate(s) is/are placed back in the 37°C incubator for three days. After three days 10 ml of stock alamar blue (Biosource Cat# DAL1100) is added to each well and the plate(s) is/are placed back in the 37°C incubator for four hours. The plate(s) are then read at 530nm

excitation and 590nm emission using the CytoFluor fluorescence reader. Direct output is recorded in relative fluorescence units.

[1215] Alamar blue is an oxidation-reduction indicator that both fluoresces and changes color in response to chemical reduction of growth medium resulting from cell growth. As cells grow in culture, innate metabolic activity results in a chemical reduction of the immediate surrounding environment. Reduction related to growth causes the indicator to change from oxidized (non-fluorescent blue) form to reduced (fluorescent red) form (i.e., stimulated proliferation will produce a stronger signal and inhibited proliferation will produce a weaker signal and the total signal is proportional to the total number of cells as well as their metabolic activity). The background level of activity is observed with the starvation medium alone. This is compared to the output observed from the positive control samples (bFGF in growth medium) and protein dilutions.

Example 45: Detection of Inhibition of a Mixed Lymphocyte Reaction

[1216] This assay can be used to detect and evaluate inhibition of a Mixed Lymphocyte Reaction (MLR) by gene products (e.g., isolated polypeptides). Inhibition of a MLR may be due to a direct effect on cell proliferation and viability, modulation of costimulatory molecules on interacting cells, modulation of adhesiveness between lymphocytes and accessory cells, or modulation of cytokine production by accessory cells. Multiple cells may be targeted by these polypeptides since the peripheral blood mononuclear fraction used in this assay includes T, B and natural killer lymphocytes, as well as monocytes and dendritic cells.

[1217] Polypeptides of interest found to inhibit the MLR may find application in diseases associated with lymphocyte and monocyte activation or proliferation. These include, but are not limited to, diseases such as asthma, arthritis, diabetes, inflammatory skin conditions, psoriasis, eczema, systemic lupus erythematosus, multiple sclerosis, glomerulonephritis, inflammatory bowel disease, crohn's disease, ulcerative colitis, arteriosclerosis, cirrhosis, graft vs. host disease, host vs. graft disease, hepatitis, leukemia and lymphoma.

[1218] Briefly, PBMCs from human donors are purified by density gradient centrifugation using Lymphocyte Separation Medium (LSM[®], density 1.0770 g/ml,

Organon Teknika Corporation, West Chester, PA). PBMCs from two donors are adjusted to 2×10^6 cells/ml in RPMI-1640 (Life Technologies, Grand Island, NY) supplemented with 10% FCS and 2 mM glutamine. PBMCs from a third donor is adjusted to 2×10^5 cells/ml. Fifty microliters of PBMCs from each donor is added to wells of a 96-well round bottom microtiter plate. Dilutions of test materials (50 μ l) is added in triplicate to microtiter wells. Test samples (of the protein of interest) are added for final dilution of 1:4; rhuIL-2 (R&D Systems, Minneapolis, MN, catalog number 202-IL) is added to a final concentration of 1 μ g/ml; anti-CD4 mAb (R&D Systems, clone 34930.11, catalog number MAB379) is added to a final concentration of 10 μ g/ml. Cells are cultured for 7-8 days at 37°C in 5% CO₂, and 1 μ C of [³H] thymidine is added to wells for the last 16 hrs of culture. Cells are harvested and thymidine incorporation determined using a Packard TopCount. Data is expressed as the mean and standard deviation of triplicate determinations.

[1219] Samples of the protein of interest are screened in separate experiments and compared to the negative control treatment, anti-CD4 mAb, which inhibits proliferation of lymphocytes and the positive control treatment, IL-2 (either as recombinant material or supernatant), which enhances proliferation of lymphocytes.

[1220] One skilled in the art could easily modify the exemplified studies to test the activity of polynucleotides (e.g., gene therapy), antibodies, agonists, and/or antagonists and fragments and variants thereof.

Example 46: Assays for Protease Activity

[1221] The following assay may be used to assess protease activity of the polypeptides of the invention.

[1222] Gelatin and casein zymography are performed essentially as described (Heusen et al., *Anal. Biochem.*, 102:196-202 (1980); Wilson et al., *Journal of Urology*, 149:653-658 (1993)). Samples are run on 10% polyacrylamide/0.1% SDS gels containing 1% gelatin or casein, soaked in 2.5% triton at room temperature for 1 hour, and in 0.1M glycine, pH 8.3 at 37°C 5 to 16 hours. After staining in amido black areas of proteolysis appear as clear areas against the blue-black background. Trypsin (Sigma T8642) is used as

a positive control.

[1223] Protease activity is also determined by monitoring the cleavage of n-a-benzoyl-L-arginine ethyl ester (BAEE) (Sigma B-4500. Reactions are set up in (25mMNaPO₄, 1mM EDTA, and 1mM BAEE), pH 7.5. Samples are added and the change in adsorbance at 260nm is monitored on the Beckman DU-6 spectrophotometer in the time-drive mode. Trypsin is used as a positive control.

[1224] Additional assays based upon the release of acid-soluble peptides from casein or hemoglobin measured as adsorbance at 280 nm or colorimetrically using the Folin method are performed as described in Bergmeyer, et al., *Methods of Enzymatic Analysis*, 5 (1984). Other assays involve the solubilization of chromogenic substrates (Ward, *Applied Science*, 251-317 (1983)).

Example 47: Identifying Serine Protease Substrate Specificity

[1225] Methods known in the art or described herein may be used to determine the substrate specificity of the polypeptides of the present invention having serine protease activity. A preferred method of determining substrate specificity is by the use of positional scanning synthetic combinatorial libraries as described in GB 2 324 529 (incorporated herein in its entirety).

Example 48: Ligand Binding Assays

[1226] The following assay may be used to assess ligand binding activity of the polypeptides of the invention.

[1227] Ligand binding assays provide a direct method for ascertaining receptor pharmacology and are adaptable to a high throughput format. The purified ligand for a polypeptide is radiolabeled to high specific activity (50-2000 Ci/mmol) for binding studies. A determination is then made that the process of radiolabeling does not diminish the activity of the ligand towards its polypeptide. Assay conditions for buffers, ions, pH and other modulators such as nucleotides are optimized to establish a workable signal to noise ratio for both membrane and whole cell polypeptide sources. For these assays, specific polypeptide binding is defined as total associated radioactivity minus the

radioactivity measured in the presence of an excess of unlabeled competing ligand. Where possible, more than one competing ligand is used to define residual nonspecific binding.

Example 49: Functional Assay in *Xenopus* Oocytes

[1228] Capped RNA transcripts from linearized plasmid templates encoding the polypeptides of the invention are synthesized in vitro with RNA polymerases in accordance with standard procedures. In vitro transcripts are suspended in water at a final concentration of 0.2 mg/ml. Ovarian lobes are removed from adult female toads, Stage V defolliculated oocytes are obtained, and RNA transcripts (10 ng/oocyte) are injected in a 50 nl bolus using a microinjection apparatus. Two electrode voltage clamps are used to measure the currents from individual *Xenopus oocytes* in response polypeptides and polypeptide agonist exposure. Recordings are made in Ca²⁺ free Barth's medium at room temperature. The *Xenopus* system can be used to screen known ligands and tissue/cell extracts for activating ligands.

Example 50: Microphysiometric Assays

[1229] Activation of a wide variety of secondary messenger systems results in extrusion of small amounts of acid from a cell. The acid formed is largely as a result of the increased metabolic activity required to fuel the intracellular signaling process. The pH changes in the media surrounding the cell are very small but are detectable by the CYTOSENSOR microphysiometer (Molecular Devices Ltd., Menlo Park, Calif.). The CYTOSENSOR is thus capable of detecting the activation of polypeptide which is coupled to an energy utilizing intracellular signaling pathway.

Example 51: Extract/Cell Supernatant Screening

[1230] A large number of mammalian receptors exist for which there remains, as yet, no cognate activating ligand (agonist). Thus, active ligands for these receptors may not be included within the ligands banks as identified to date. Accordingly, the polypeptides of the invention can also be functionally screened (using calcium, cAMP, microphysiometer,

oocyte electrophysiology, etc., functional screens) against tissue extracts to identify its natural ligands. Extracts that produce positive functional responses can be sequentially subfractionated until an activating ligand is isolated and identified.

Example 52: Calcium and cAMP Functional Assays

[1231] Seven transmembrane receptors which are expressed in HEK 293 cells have been shown to be coupled functionally to activation of PLC and calcium mobilization and/or cAMP stimulation or inhibition. Basal calcium levels in the HEK 293 cells in receptor-transfected or vector control cells were observed to be in the normal, 100 nM to 200 nM, range. HEK 293 cells expressing recombinant receptors are loaded with fura 2 and in a single day >150 selected ligands or tissue/cell extracts are evaluated for agonist induced calcium mobilization. Similarly, HEK 293 cells expressing recombinant receptors are evaluated for the stimulation or inhibition of cAMP production using standard cAMP quantitation assays. Agonists presenting a calcium transient or cAMP fluctuation are tested in vector control cells to determine if the response is unique to the transfected cells expressing receptor.

Example 53: ATP-binding assay

[1232] The following assay may be used to assess ATP-binding activity of polypeptides of the invention.

[1233] ATP-binding activity of the polypeptides of the invention may be detected using the ATP-binding assay described in U.S. Patent 5,858,719, which is herein incorporated by reference in its entirety. Briefly, ATP-binding to polypeptides of the invention is measured via photoaffinity labeling with 8-azido-ATP in a competition assay. Reaction mixtures containing 1 mg/ml of the ABC transport protein of the present invention are incubated with varying concentrations of ATP, or the non-hydrolyzable ATP analog adenylyl-5'-imidodiphosphate for 10 minutes at 4°C. A mixture of 8-azido-ATP (Sigma Chem. Corp., St. Louis, MO.) plus 8-azido-ATP (³²P-ATP) (5 mCi/μmol, ICN, Irvine CA.) is added to a final concentration of 100 μM and 0.5 ml aliquots are placed in

the wells of a porcelain spot plate on ice. The plate is irradiated using a short wave 254 nm UV lamp at a distance of 2.5 cm from the plate for two one-minute intervals with a one-minute cooling interval in between. The reaction is stopped by addition of dithiothreitol to a final concentration of 2mM. The incubations are subjected to SDS-PAGE electrophoresis, dried, and autoradiographed. Protein bands corresponding to the particular polypeptides of the invention are excised, and the radioactivity quantified. A decrease in radioactivity with increasing ATP or adenyly-5'-imidodiphosphate provides a measure of ATP affinity to the polypeptides.

Example 54: Small Molecule Screening

[1234] This invention is particularly useful for screening therapeutic compounds by using the polypeptides of the invention, or binding fragments thereof, in any of a variety of drug screening techniques. The polypeptide or fragment employed in such a test may be affixed to a solid support, expressed on a cell surface, free in solution, or located intracellularly. One method of drug screening utilizes eukaryotic or prokaryotic host cells which are stably transformed with recombinant nucleic acids expressing the polypeptide or fragment. Drugs are screened against such transformed cells in competitive binding assays. One may measure, for example, the formulation of complexes between the agent being tested and polypeptide of the invention.

[1235] Thus, the present invention provides methods of screening for drugs or any other agents which affect activities mediated by the polypeptides of the invention. These methods comprise contacting such an agent with a polypeptide of the invention or fragment thereof and assaying for the presence of a complex between the agent and the polypeptide or fragment thereof, by methods well known in the art. In such a competitive binding assay, the agents to screen are typically labeled. Following incubation, free agent is separated from that present in bound form, and the amount of free or uncomplexed label is a measure of the ability of a particular agent to bind to the polypeptides of the invention.

[1236] Another technique for drug screening provides high throughput screening for compounds having suitable binding affinity to the polypeptides of the invention, and is described in great detail in European Patent Application 84/03564, published on

September 13, 1984, which is herein incorporated by reference in its entirety. Briefly stated, large numbers of different small molecule test compounds are synthesized on a solid substrate, such as plastic pins or some other surface. The test compounds are reacted with polypeptides of the invention and washed. Bound polypeptides are then detected by methods well known in the art. Purified polypeptides are coated directly onto plates for use in the aforementioned drug screening techniques. In addition, non-neutralizing antibodies may be used to capture the peptide and immobilize it on the solid support.

[1237] This invention also contemplates the use of competitive drug screening assays in which neutralizing antibodies capable of binding polypeptides of the invention specifically compete with a test compound for binding to the polypeptides or fragments thereof. In this manner, the antibodies are used to detect the presence of any peptide which shares one or more antigenic epitopes with a polypeptide of the invention.

Example 55: Phosphorylation Assay

[1238] In order to assay for phosphorylation activity of the polypeptides of the invention, a phosphorylation assay as described in U.S. Patent 5,958,405 (which is herein incorporated by reference) is utilized. Briefly, phosphorylation activity may be measured by phosphorylation of a protein substrate using gamma-labeled ^{32}P -ATP and quantitation of the incorporated radioactivity using a gamma radioisotope counter. The polypeptides of the invention are incubated with the protein substrate, ^{32}P -ATP, and a kinase buffer. The ^{32}P incorporated into the substrate is then separated from free ^{32}P -ATP by electrophoresis, and the incorporated ^{32}P is counted and compared to a negative control. Radioactivity counts above the negative control are indicative of phosphorylation activity of the polypeptides of the invention.

Example 56: Detection of Phosphorylation Activity (Activation) of the Polypeptides of the Invention in the Presence of Polypeptide Ligands

[1239] Methods known in the art or described herein may be used to determine the phosphorylation activity of the polypeptides of the invention. A preferred method of

determining phosphorylation activity is by the use of the tyrosine phosphorylation assay as described in US 5,817,471 (incorporated herein by reference).

***Example 57: Identification Of Signal Transduction Proteins That Interact
With Polypeptides Of The Present Invention***

[1240] The purified polypeptides of the invention are research tools for the identification, characterization and purification of additional signal transduction pathway proteins or receptor proteins. Briefly, labeled polypeptides of the invention are useful as reagents for the purification of molecules with which it interacts. In one embodiment of affinity purification, polypeptides of the invention are covalently coupled to a chromatography column. Cell-free extract derived from putative target cells, such as carcinoma tissues, is passed over the column, and molecules with appropriate affinity bind to the polypeptides of the invention. The protein complex is recovered from the column, dissociated, and the recovered molecule subjected to N-terminal protein sequencing. This amino acid sequence is then used to identify the captured molecule or to design degenerate oligonucleotide probes for cloning the relevant gene from an appropriate cDNA library.

Example 58: IL-6 Bioassay

[1241] To test the proliferative effects of the polypeptides of the invention, the IL-6 Bioassay as described by Marz *et al.* is utilized (*Proc. Natl. Acad. Sci., U.S.A.*, 95:3251-56 (1998), which is herein incorporated by reference). Briefly, IL-6 dependent B9 murine cells are washed three times in IL-6 free medium and plated at a concentration of 5,000 cells per well in 50 μ l, and 50 μ l of the IL-6-like polypeptide is added. After 68 hrs. at 37°C, the number of viable cells is measured by adding the tetrazolium salt thiazolyl blue (MTT) and incubating for a further 4 hrs. at 37°C. B9 cells are lysed by SDS and optical density is measured at 570 nm. Controls containing IL-6 (positive) and no cytokine (negative) are utilized. Enhanced proliferation in the test sample(s) relative to the negative control is indicative of proliferative effects mediated by polypeptides of the invention.

Example 59: Support of Chicken Embryo Neuron Survival

[1242] To test whether sympathetic neuronal cell viability is supported by polypeptides of the invention, the chicken embryo neuronal survival assay of Senaldi *et al* is utilized (*Proc. Natl. Acad. Sci., U.S.A.*, 96:11458-63 (1998), which is herein incorporated by reference). Briefly, motor and sympathetic neurons are isolated from chicken embryos, resuspended in L15 medium (with 10% FCS, glucose, sodium selenite, progesterone, conalbumin, putrescine, and insulin; Life Technologies, Rockville, MD.) and Dulbecco's modified Eagles medium [with 10% FCS, glutamine, penicillin, and 25 mM Hepes buffer (pH 7.2); Life Technologies, Rockville, MD.], respectively, and incubated at 37°C in 5% CO₂ in the presence of different concentrations of the purified IL-6-like polypeptide, as well as a negative control lacking any cytokine. After 3 days, neuron survival is determined by evaluation of cellular morphology, and through the use of the colorimetric assay of Mosmann (Mosmann, T., *J. Immunol. Methods*, 65:55-63 (1983)). Enhanced neuronal cell viability as compared to the controls lacking cytokine is indicative of the ability of the inventive purified IL-6-like polypeptide(s) to enhance the survival of neuronal cells.

Example 60: Assay for Phosphatase Activity

[1243] The following assay may be used to assess serine/threonine phosphatase (PTPase) activity of the polypeptides of the invention.

[1244] In order to assay for serine/threonine phosphatase (PTPase) activity, assays can be utilized which are widely known to those skilled in the art. For example, the serine/threonine phosphatase (PSPase) activity is measured using a PSPase assay kit from New England Biolabs, Inc. Myelin basic protein (MyBP), a substrate for PSPase, is phosphorylated on serine and threonine residues with cAMP-dependent Protein Kinase in the presence of [³²P]ATP. Protein serine/threonine phosphatase activity is then determined by measuring the release of inorganic phosphate from 32P-labeled MyBP.

Example 61: Interaction of Serine/Threonine Phosphatases with other Proteins

[1245] The polypeptides of the invention with serine/threonine phosphatase activity as determined in Example 60 are research tools for the identification, characterization and purification of additional interacting proteins or receptor proteins, or other signal transduction pathway proteins. Briefly, labeled polypeptide(s) of the invention is useful as a reagent for the purification of molecules with which it interacts. In one embodiment of affinity purification, polypeptide of the invention is covalently coupled to a chromatography column. Cell-free extract derived from putative target cells, such as neural or liver cells, is passed over the column, and molecules with appropriate affinity bind to the polypeptides of the invention. The polypeptides of the invention -complex is recovered from the column, dissociated, and the recovered molecule subjected to N-terminal protein sequencing. This amino acid sequence is then used to identify the captured molecule or to design degenerate oligonucleotide probes for cloning the relevant gene from an appropriate cDNA library.

Example 62: Assaying for Heparanase Activity

[1246] In order to assay for heparanase activity of the polypeptides of the invention, the heparanase assay described by Vlodavsky et al is utilized (Vlodavsky, I., et al., Nat. Med., 5:793-802 (1999)). Briefly, cell lysates, conditioned media or intact cells (1×10^6 cells per 35-mm dish) are incubated for 18 hrs at 37°C, pH 6.2-6.6, with ^{35}S -labeled ECM or soluble ECM derived peak I proteoglycans. The incubation medium is centrifuged and the supernatant is analyzed by gel filtration on a Sepharose CL-6B column (0.9 x 30 cm). Fractions are eluted with PBS and their radioactivity is measured. Degradation fragments of heparan sulfate side chains are eluted from Sepharose 6B at $0.5 < K_{av} < 0.8$ (peak II). Each experiment is done at least three times. Degradation fragments corresponding to "peak II," as described by Vlodavsky et al., is indicative of the activity of the polypeptides of the invention in cleaving heparan sulfate.

Example 63: Immobilization of biomolecules

[1247] This example provides a method for the stabilization of polypeptides of the invention in non-host cell lipid bilayer constructs (see, e.g., Bieri et al., Nature Biotech 17:1105-1108 (1999), hereby incorporated by reference in its entirety herein) which can be adapted for the study of polypeptides of the invention in the various functional assays described above. Briefly, carbohydrate-specific chemistry for biotinylation is used to confine a biotin tag to the extracellular domain of the polypeptides of the invention, thus allowing uniform orientation upon immobilization. A 50uM solution of polypeptides of the invention in washed membranes is incubated with 20 mM NaIO₄ and 1.5 mg/ml (4mM) BACH or 2 mg/ml (7.5mM) biotin-hydrazide for 1 hr at room temperature (reaction volume, 150ul). Then the sample is dialyzed (Pierce Slidealizer Cassett, 10 kDa cutoff; Pierce Chemical Co., Rockford IL) at 4C first for 5 h, exchanging the buffer after each hour, and finally for 12 h against 500 ml buffer R (0.15 M NaCl, 1 mM MgCl₂, 10 mM sodium phosphate, pH7). Just before addition into a cuvette, the sample is diluted 1:5 in buffer ROG50 (Buffer R supplemented with 50 mM octylglucoside).

Example 64: TAQMAN

[1248] Quantitative PCR (QPCR). Total RNA from cells in culture are extracted by Trizol separation as recommended by the supplier (LifeTechnologies). (Total RNA is treated with DNase I (Life Technologies) to remove any contaminating genomic DNA before reverse transcription.) Total RNA (50 ng) is used in a one-step, 50ul, RT-QPCR, consisting of Taqman Buffer A (Perkin-Elmer; 50 mM KCl/10 mM Tris, pH 8.3), 5.5 mM MgCl₂, 240 μ M each dNTP, 0.4 units RNase inhibitor(Promega), 8%glycerol, 0.012% Tween-20, 0.05% gelatin, 0.3uM primers, 0.1uM probe, 0.025units Amplitaq Gold (Perkin-Elmer) and 2.5 units Superscript II reverse transcriptase (Life Technologies). As a control for genomic contamination, parallel reactions are setup without reverse transcriptase. The relative abundance of (unknown) and 18S RNAs are assessed by using the Applied Biosystems Prism 7700 Sequence Detection System (Livak, K. J., Flood, S. J., Marmaro, J., Giusti, W. & Deetz, K. (1995) PCR Methods Appl. 4, 357-362).

Reactions are carried out at 48°C for 30 min, 95°C for 10 min, followed by 40 cycles of 95°C for 15s, 60°C for 1 min. Reactions are performed in triplicate.

[1249] Primers (f & r) and FRET probes sets are designed using Primer Express Software (Perkin-Elmer). Probes are labeled at the 5'-end with the reporter dye 6-FAM and on the 3'-end with the quencher dye TAMRA (Biosource International, Camarillo, CA or Perkin-Elmer).

Example 65: Assays for Metalloproteinase Activity

[1250] Metalloproteinases (EC 3.4.24.-) are peptide hydrolases which use metal ions, such as Zn^{2+} , as the catalytic mechanism. Metalloproteinase activity of polypeptides of the present invention can be assayed according to the following methods.

Proteolysis of alpha-2-macroglobulin

[1251] To confirm protease activity, purified polypeptides of the invention are mixed with the substrate alpha-2-macroglobulin (0.2 unit/ml; Boehringer Mannheim, Germany) in 1x assay buffer (50 mM HEPES, pH 7.5, 0.2 M NaCl, 10 mM $CaCl_2$, 25 μM $ZnCl_2$ and 0.05% Brij-35) and incubated at 37°C for 1-5 days. Trypsin is used as positive control. Negative controls contain only alpha-2-macroglobulin in assay buffer. The samples are collected and boiled in SDS-PAGE sample buffer containing 5% 2-mercaptoethanol for 5-min, then loaded onto 8% SDS-polyacrylamide gel. After electrophoresis the proteins are visualized by silver staining. Proteolysis is evident by the appearance of lower molecular weight bands as compared to the negative control.

Inhibition of alpha-2-macroglobulin proteolysis by inhibitors of metalloproteinases

[1252] Known metalloproteinase inhibitors (metal chelators (EDTA, EGTA, AND $HgCl_2$), peptide metalloproteinase inhibitors (TIMP-1 and TIMP-2), and commercial small molecule MMP inhibitors) are used to characterize the proteolytic activity of polypeptides of the invention. The three synthetic MMP inhibitors used are: MMP inhibitor I, [IC_{50} = 1.0 μM against MMP-1 and MMP-8; IC_{50} = 30 μM against MMP-9; IC_{50} = 150 μM against MMP-3]; MMP-3 (stromelysin-1) inhibitor I [IC_{50} = 5 μM against MMP-3], and MMP-3 inhibitor II [K_i = 130 nM against MMP-3]; inhibitors available

through Calbiochem, catalog # 444250, 444218, and 444225, respectively). Briefly, different concentrations of the small molecule MMP inhibitors are mixed with purified polypeptides of the invention (50 μ g/ml) in 22.9 μ l of 1x HEPES buffer (50 mM HEPES, pH 7.5, 0.2 M NaCl, 10 mM CaCl₂, 25 μ M ZnCl₂ and 0.05%Brij-35) and incubated at room temperature (24 °C) for 2-hr, then 7.1 μ l of substrate alpha-2-macroglobulin (0.2 unit/ml) is added and incubated at 37°C for 20-hr. The reactions are stopped by adding 4x sample buffer and boiled immediately for 5 minutes. After SDS-PAGE, the protein bands are visualized by silver stain.

Synthetic Fluorogenic Peptide Substrates Cleavage Assay

[1253] The substrate specificity for polypeptides of the invention with demonstrated metalloproteinase activity can be determined using synthetic fluorogenic peptide substrates (purchased from BACHEM Bioscience Inc). Test substrates include, M-1985, M-2225, M-2105, M-2110, and M-2255. The first four are MMP substrates and the last one is a substrate of tumor necrosis factor- α (TNF- α) converting enzyme (TACE). All the substrates are prepared in 1:1 dimethyl sulfoxide (DMSO) and water. The stock solutions are 50-500 μ M. Fluorescent assays are performed by using a Perkin Elmer LS 50B luminescence spectrometer equipped with a constant temperature water bath. The excitation λ is 328 nm and the emission λ is 393 nm. Briefly, the assay is carried out by incubating 176 μ l 1x HEPES buffer (0.2 M NaCl, 10 mM CaCl₂, 0.05% Brij-35 and 50 mM HEPES, pH 7.5) with 4 μ l of substrate solution (50 μ M) at 25 °C for 15 minutes, and then adding 20 μ l of a purified polypeptide of the invention into the assay cuvet. The final concentration of substrate is 1 μ M. Initial hydrolysis rates are monitored for 30-min.

Example 66: Characterization of the cDNA contained in a deposited plasmid

[1254] The size of the cDNA insert contained in a deposited plasmid may be routinely determined using techniques known in the art, such as PCR amplification using synthetic primers hybridizable to the 3' and 5' ends of the cDNA sequence. For example, two

primers of 17-30 nucleotides derived from each end of the cDNA (i.e., hybridizable to the absolute 5' nucleotide or the 3' nucleotide end of the sequence of SEQ ID NO:X, respectively) are synthesized and used to amplify the cDNA using the deposited cDNA plasmid as a template. The polymerase chain reaction is carried out under routine conditions, for instance, in 25 ul of reaction mixture with 0.5 ug of the above cDNA template. A convenient reaction mixture is 1.5-5 mM MgCl₂, 0.01% (w/v) gelatin, 20 uM each of dATP, dCTP, dGTP, dTTP, 25 pmol of each primer and 0.25 Unit of Taq polymerase. Thirty five cycles of PCR (denaturation at 94 degree C for 1 min; annealing at 55 degree C for 1 min; elongation at 72 degree C for 1 min) are performed with a Perkin-Elmer Cetus automated thermal cycler. The amplified product is analyzed by agarose gel electrophoresis. The PCR product is verified to be the selected sequence by subcloning and sequencing the DNA product.

[1255] Use of the above methodologies and/or other methodologies known in the art can be used to generate fragments from the clone.

[1256] It will be clear that the invention may be practiced otherwise than as particularly described in the foregoing description and examples. Numerous modifications and variations of the present invention are possible in light of the above teachings and, therefore, are within the scope of the appended claims.

[1257] The entire disclosure of each document cited (including patents, patent applications, journal articles, abstracts, laboratory manuals, books, or other disclosures) in the Background of the Invention, Detailed Description, and Examples is hereby incorporated herein by reference. In addition, the CD-R copy of the sequence listing submitted herewith and the corresponding computer readable form are both incorporated herein by reference in their entireties. The specification and sequence listing of each of the following U.S. applications are herein incorporated by reference in their entirety: Application No. 60/179,065, filed on 31-Jan-2000; Application No. 60/180,628, filed on 04-Feb-2000; and Application No. 60/235,484, filed on 26-Sep-2000.

[1258] Moreover, the microfiche copy and the corresponding computer readable form of the Sequence Listing of U.S. Application Serial No. 60/179,065, and the hard copy of and the corresponding computer readable form of the Sequence Listing of U.S.

Application Serial No. 60/180,628 and 60/235,484 are also incorporated herein by reference in their entireties.

INDICATIONS RELATING TO A DEPOSITED MICROORGANISM OR OTHER BIOLOGICAL MATERIAL (PCT Rule 13bis)			
A. The indications made below relate to the deposited microorganism or other biological material referred to in the description on page 16, line 2.			
B. IDENTIFICATION OF DEPOSIT		Further deposits are identified on an additional sheet <input checked="" type="checkbox"/>	
Name of depositary institution: American Type Culture Collection			
Address of depositary institution <i>(including postal code and country)</i> 10801 University Boulevard Manassas, Virginia 20110-2209 United States of America			
Date of deposit 12 September 2000		Accession Number PTA-2448	
C. ADDITIONAL INDICATIONS <i>(leave blank if not applicable)</i>		This information is continued on an additional sheet <input type="checkbox"/>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE <i>(if the indications are not for all designated States)</i>			
Europe In respect of those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which the application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28(4) EPC). <div style="text-align: right;">Continued on additional sheets</div>			
E. SEPARATE FURNISHING OF INDICATIONS <i>(leave blank if not applicable)</i>			
The indications listed below will be submitted to the international Bureau later <i>(specify the general nature of the indications e.g., "Accession Number of Deposit")</i>			
For receiving Office use only		For International Bureau use only	
<input type="checkbox"/> This sheet was received with the international application		<input type="checkbox"/> This sheet was received by the International Bureau on:	
Authorized officer		Authorized officer	

ATCC Deposit No. PTA-2448

CANADA

The applicant requests that, until either a Canadian patent has been issued on the basis of an application or the application has been refused, or is abandoned and no longer subject to reinstatement, or is withdrawn, the Commissioner of Patents only authorizes the furnishing of a sample of the deposited biological material referred to in the application to an independent expert nominated by the Commissioner, the applicant must, by a written statement, inform the International Bureau accordingly before completion of technical preparations for publication of the international application.

NORWAY

The applicant hereby requests that the application has been laid open to public inspection (by the Norwegian Patent Office), or has been finally decided upon by the Norwegian Patent Office without having been laid open inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the Norwegian Patent Office not later than at the time when the application is made available to the public under Sections 22 and 33(3) of the Norwegian Patents Act. If such a request has been filed by the applicant, any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on the list of recognized experts drawn up by the Norwegian Patent Office or any person approved by the applicant in the individual case.

AUSTRALIA

The applicant hereby gives notice that the furnishing of a sample of a microorganism shall only be effected prior to the grant of a patent, or prior to the lapsing, refusal or withdrawal of the application, to a person who is a skilled addressee without an interest in the invention (Regulation 3.25(3) of the Australian Patents Regulations).

FINLAND

The applicant hereby requests that, until the application has been laid open to public inspection (by the National Board of Patents and Regulations), or has been finally decided upon by the National Board of Patents and Registration without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art.

ATCC Deposit No.: PTA-2448

UNITED KINGDOM

The applicant hereby requests that the furnishing of a sample of a microorganism shall only be made available to an expert. The request to this effect must be filed by the applicant with the International Bureau before the completion of the technical preparations for the international publication of the application.

DENMARK

The applicant hereby requests that, until the application has been laid open to public inspection (by the Danish Patent Office), or has been finally decided upon by the Danish Patent office without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the Danish Patent Office not later than at the time when the application is made available to the public under Sections 22 and 33(3) of the Danish Patents Act. If such a request has been filed by the applicant, any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on a list of recognized experts drawn up by the Danish Patent Office or any person by the applicant in the individual case.

SWEDEN

The applicant hereby requests that, until the application has been laid open to public inspection (by the Swedish Patent Office), or has been finally decided upon by the Swedish Patent Office without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the International Bureau before the expiration of 16 months from the priority date (preferably on the Form PCT/RO/134 reproduced in annex Z of Volume I of the PCT Applicant's Guide). If such a request has been filed by the applicant any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on a list of recognized experts drawn up by the Swedish Patent Office or any person approved by a applicant in the individual case.

NETHERLANDS

The applicant hereby requests that until the date of a grant of a Netherlands patent or until the date on which the application is refused or withdrawn or lapsed, the microorganism shall be made available as provided in the 31F(1) of the Patent Rules only by the issue of a sample to an expert. The request to this effect must be furnished by the applicant with the Netherlands Industrial Property Office before the date on which the application is made available to the public under Section 22C or Section 25 of the Patents Act of the Kingdom of the Netherlands, whichever of the two dates occurs earlier.

**INDICATIONS RELATING TO A DEPOSITED MICROORGANISM
OR OTHER BIOLOGICAL MATERIAL**

(PCT Rule 13bis)

A. The indications made below relate to the deposited microorganism or other biological material referred to in the description on page 16, line 3.

B. IDENTIFICATION OF DEPOSIT

Further deposits are identified on an additional sheet ☒

Name of depositary institution: American Type Culture Collection

Address of depositary institution (including postal code and country)

10801 University Boulevard
Manassas, Virginia 20110-2209
United States of America

Date of deposit
12 September 2000

Accession Number
PTA-2449

C. ADDITIONAL INDICATIONS (leave blank if not applicable)

This information is continued on an additional sheet ☐

D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)

Europe

In respect of those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which the application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28(4) EPC).
Continued on additional sheets

E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)

The indications listed below will be submitted to the international Bureau later (specify the general nature of the indications e.g., "Accession Number of Deposit")

For receiving Office use only				For International Bureau use only	
<input checked="" type="checkbox"/> This sheet was received with the international application			<input type="checkbox"/> This sheet was received by the International Bureau on:		
Authorized officer MELVIN S. BRIGGS SR. INTERNATIONAL DIVISION 703-305-5163 <i>MB</i>			Authorized officer		

ATCC Deposit No. PTA-2449**CANADA**

The applicant requests that, until either a Canadian patent has been issued on the basis of an application or the application has been refused, or is abandoned and no longer subject to reinstatement, or is withdrawn, the Commissioner of Patents only authorizes the furnishing of a sample of the deposited biological material referred to in the application to an independent expert nominated by the Commissioner, the applicant must, by a written statement, inform the International Bureau accordingly before completion of technical preparations for publication of the international application.

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FINLAND

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ATCC Deposit No.: PTA-2449

UNITED KINGDOM

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What Is Claimed Is:

1. An isolated nucleic acid molecule comprising a polynucleotide having a nucleotide sequence at least 95% identical to a sequence selected from the group consisting of:
 - (a) a polynucleotide fragment of SEQ ID NO:X or a polynucleotide fragment of the cDNA sequence contained in Clone ID NO:Z, which is hybridizable to SEQ ID NO:X;
 - (b) a polynucleotide encoding a polypeptide fragment of SEQ ID NO:Y or a polypeptide fragment encoded by the cDNA sequence contained in cDNA Clone ID NO:Z, which is hybridizable to SEQ ID NO:X;
 - (c) a polynucleotide encoding a polypeptide fragment of a polypeptide encoded by SEQ ID NO:X or a polypeptide fragment encoded by the cDNA sequence contained in cDNA Clone ID NO:Z, which is hybridizable to SEQ ID NO:X;
 - (d) a polynucleotide encoding a polypeptide domain of SEQ ID NO:Y or a polypeptide domain encoded by the cDNA sequence contained in cDNA Clone ID NO:Z, which is hybridizable to SEQ ID NO:X;
 - (e) a polynucleotide encoding a polypeptide epitope of SEQ ID NO:Y or a polypeptide epitope encoded by the cDNA sequence contained in cDNA Clone ID NO:Z, which is hybridizable to SEQ ID NO:X;
 - (f) a polynucleotide encoding a polypeptide of SEQ ID NO:Y or the cDNA sequence contained in cDNA Clone ID NO:Z, which is hybridizable to SEQ ID NO:X, having biological activity;
 - (g) a polynucleotide which is a variant of SEQ ID NO:X;
 - (h) a polynucleotide which is an allelic variant of SEQ ID NO:X;
 - (i) a polynucleotide which encodes a species homologue of the SEQ ID NO:Y;
 - (j) a polynucleotide capable of hybridizing under stringent conditions to any one of the polynucleotides specified in (a)-(i), wherein said polynucleotide does not hybridize under stringent conditions to a nucleic acid molecule having a nucleotide sequence of only A residues or of only T residues.
2. The isolated nucleic acid molecule of claim 1, wherein the polynucleotide fragment comprises a nucleotide sequence encoding a protein.

3. The isolated nucleic acid molecule of claim 1, wherein the polynucleotide fragment comprises a nucleotide sequence encoding the sequence identified as SEQ ID NO:Y or the polypeptide encoded by the cDNA sequence contained in cDNA Clone ID NO:Z, which is hybridizable to SEQ ID NO:X.
4. The isolated nucleic acid molecule of claim 1, wherein the polynucleotide fragment comprises the entire nucleotide sequence of SEQ ID NO:X or the cDNA sequence contained in cDNA Clone ID NO:Z, which is hybridizable to SEQ ID NO:X.
5. The isolated nucleic acid molecule of claim 2, wherein the nucleotide sequence comprises sequential nucleotide deletions from either the C-terminus or the N-terminus.
6. The isolated nucleic acid molecule of claim 3, wherein the nucleotide sequence comprises sequential nucleotide deletions from either the C-terminus or the N-terminus.
7. A recombinant vector comprising the isolated nucleic acid molecule of claim 1.
8. A method of making a recombinant host cell comprising the isolated nucleic acid molecule of claim 1.
9. A recombinant host cell produced by the method of claim 8.
10. The recombinant host cell of claim 9 comprising vector sequences.
11. An isolated polypeptide comprising an amino acid sequence at least 90% identical to a sequence selected from the group consisting of:
 - (a) a polypeptide fragment of SEQ ID NO:Y or the encoded sequence contained in cDNA Clone ID NO:Z;

(b) a polypeptide fragment of SEQ ID NO:Y or the encoded sequence contained in cDNA Clone ID NO:Z, having biological activity;

(c) a polypeptide domain of SEQ ID NO:Y or the encoded sequence contained in cDNA Clone ID NO:Z;

(d) a polypeptide epitope of SEQ ID NO:Y or the encoded sequence contained in cDNA Clone ID NO:Z;

(e) a full length protein of SEQ ID NO:Y or the encoded sequence contained in cDNA Clone ID NO:Z;

(f) a variant of SEQ ID NO:Y;

(g) an allelic variant of SEQ ID NO:Y; or

(h) a species homologue of the SEQ ID NO:Y.

12. The isolated polypeptide of claim 11, wherein the full length protein comprises sequential amino acid deletions from either the C-terminus or the N-terminus.

13. An isolated antibody that binds specifically to the isolated polypeptide of claim 11.

14. A recombinant host cell that expresses the isolated polypeptide of claim 11.

15. A method of making an isolated polypeptide comprising:

(a) culturing the recombinant host cell of claim 14 under conditions such that said polypeptide is expressed; and

(b) recovering said polypeptide.

16. The polypeptide produced by claim 15.

17. A method for preventing, treating, or ameliorating a medical condition, comprising administering to a mammalian subject a therapeutically effective amount of the polynucleotide of claim 1.

18. A method of diagnosing a pathological condition or a susceptibility to a pathological condition in a subject comprising:

- (a) determining the presence or absence of a mutation in the polynucleotide of claim 1; and
- (b) diagnosing a pathological condition or a susceptibility to a pathological condition based on the presence or absence of said mutation.

19. A method of diagnosing a pathological condition or a susceptibility to a pathological condition in a subject comprising:

- (a) determining the presence or amount of expression of the polypeptide of claim 11 in a biological sample; and
- (b) diagnosing a pathological condition or a susceptibility to a pathological condition based on the presence or amount of expression of the polypeptide.

20. A method for identifying a binding partner to the polypeptide of claim 11 comprising:

- (a) contacting the polypeptide of claim 11 with a binding partner; and
- (b) determining whether the binding partner effects an activity of the polypeptide.

21. The gene corresponding to the cDNA sequence of SEQ ID NO:Y.

22. A method of identifying an activity in a biological assay, wherein the method comprises:

- (a) expressing SEQ ID NO:X in a cell;
- (b) isolating the supernatant;
- (c) detecting an activity in a biological assay; and
- (d) identifying the protein in the supernatant having the activity.

23. The product produced by the method of claim 20.

24. A method for preventing, treating, or ameliorating a medical condition, comprising administering to a mammalian subject a therapeutically effective amount of the polypeptide of claim 11.